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(54) Title: THERAPEUTIC USE OF AN INHIBITOR OF A HEDGEHOG OR A HEDGEHOG-RELATED SIGNALLING PATHWAY

(57) Abstract: Use of an inhibitor of a Hedgehog signalling pathway, or an inhibitor of a pathway which is a target of the Hedgehog signalling pathway in the preparation of a medicament for treatment of epithelial cell hyperplasia, fibrosis of tissue, inflammation, cancer or an immune disorder. Also a transgenic animal or cell line capable of expressing a component or an inhibitor of a hedgehog signalling pathway or a target pathway of the hedgehog signalling pathway.

Therapeutic Use

Field of the Invention

- 5 The present invention relates to a novel therapeutic use, a composition for use in such therapy, a method for identifying useful compounds, a vector and a transgenic animal capable of expressing such compounds.

Background

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In many tissues, such as the lung and kidney, chronic unresolving inflammation may lead to remodelling in which both epithelial cell hyperplasia and fibrosis occur. In addition, there is an accompanying mononuclear cell infiltration at local sites of inflammation and the induction of immune responses reactive with self antigens. Similar pathology is also observed in chronic rejection of transplanted organs. In general, these diseases are difficult to manage clinically and this is well illustrated by chronic obstructive pulmonary disease where current pharmacological intervention has limited effects. Therefore, the ability to control this dysregulation of epithelial repair processes that drive anti-self immune responses and tissue remodelling will have an important clinical impact.

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Summary of the Invention

The formation of epithelial surfaces and the regulation of epithelial cell growth appears to be highly conserved among species. Recent studies in developmental biology have demonstrated the importance of epithelial cell growth factor genes Shh and Wnt 1 and the TGF- β superfamily members BMP4 and BMP7. These gene products not only play an important role in developmental epithelial cell growth but also in tissue patterning. The latter requires the coordinated and

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temporal regulation of cell differentiation programmes. During development cells respond to growth factors which are found within their environment and it is the interpretation of these various growth factor signalling pathways that will determine the differentiation programme that is initiated. To date little is known
5 of the contribution of these genes in tissue remodelling and fibrosis that are observed in chronic inflammation; although BMP has been implicated in fibrosis in bone disease. We have noted that in diseased lungs there is increased expression of the gene *patched* which is involved in Hedgehog signalling. We have also noted that intratracheal instillation of plasmid DNA for the Hedgehog
10 gene leads to the development of epithelial cell hyperplasia, fibrosis of tissue and the infiltration of mononuclear cells. The pathology is similar to that observed in interstitial lung disease.

We propose that antagonists of components of Hedgehog signalling and/or
15 antagonists of components of a signalling pathway which is a target of Hedgehog signalling may prevent and/or reverse diseases such as epithelial cell hyperplasia, tissue fibrosis, chronic inflammation, cancer and also prevent graft rejection.

20 Statements of Invention

In one aspect the present invention provides a method of treatment for epithelial cell hyperplasia, fibrosis of tissue, inflammation, cancer, or an immune disorder comprising the administration of a therapeutically effective amount of an
25 antagonist of a component of a Hedgehog family member signalling pathway or an antagonist of a component of a signalling pathway which is a target of Hedgehog signalling to an individual in need of the same.

Put another way the present invention provides use of an antagonist of a
30 component of a Hedgehog family member signalling pathway or an antagonist of

a component of a signalling pathway which is a target of Hedgehog signalling in the preparation of a medicament for the treatment of epithelial cell hyperplasia, fibrosis of tissue, cancer, inflammation, or an immune disorder.

- 5 In one embodiment the Hedgehog family member is Sonic hedgehog, Indian hedgehog or Desert hedgehog.

In one embodiment the pathway which is a target of Hedgehog signalling is a BMP signalling pathway or a Wnt signalling pathway.

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In one embodiment the antagonist is HIP, cyclopamine, Fzb, Cerberus, WIF-1, Xnr-3, Noggin, Chordin, Gremlin, or Follistatin or a derivative, fragment, variant, mimetic, homologue or analogue thereof.

- 15 In another embodiment the antagonist is an antibody to a component of the Hedgehog signalling pathway or an antibody to a component of the target pathway of Hedgehog signalling.

20 In a further embodiment the antagonist is itself a component of the Hedgehog signalling pathway or a component of the target pathway of Hedgehog signalling.

Preferably the method of the present invention relates to the treatment of pulmonary hyperplasia or pulmonary fibrosis.

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More preferably the method of the present invention relates to the treatment of adult respiratory distress syndrome; chronic obstructive airway disorders including asthma, emphysema and chronic bronchitis; atelectasis; occupational lung disease including silicosis; hypersensitivity diseases of the lung including
30 hypersensitivity pneumonitis; idiopathic interstitial lung diseases including

idiopathic pulmonary fibrosis, pneumonia including usual interstitial pneumonia, desquamative interstitial pneumonia and acute interstitial pneumonia; and pleural fibrosis.

- 5 In one embodiment the immune disorder is an autoimmune disease or graft rejection.

More particularly, the autoimmune disease may be thyroiditis, insultitis, multiple sclerosis, iridocyclitis, uveitis, orchitis, hepatitis, Addison's disease, myasthenia
10 gravis, rheumatoid arthritis and lupus erythematosus.

In one embodiment the cancer is an adenocarcinoma.

In another aspect the present invention provides a composition for use in
15 treatment of epithelial cell hyperplasia, fibrosis of tissue, inflammation, cancer or an immune disorder comprising a therapeutically effective amount of an antagonist of a Hedgehog signalling pathway or an antagonist of the target pathway of Hedgehog signalling and a pharmaceutically acceptable carrier, diluent or excipient.

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In another aspect the present invention provides a method for identifying a compound that is an inhibitor of a Hedgehog signalling pathway or a target pathway of the Hedgehog signalling pathway comprising the steps of: (a) determining the activity of the signalling pathway in the presence and absence of
25 said compound; (b) comparing the activities observed in step (a); and (c) identifying said compound as inhibitor by the observed difference in the activity of the pathway in the presence and absence of said compound.

In yet another aspect the present invention provides a vector capable of expressing an antagonist of a component of a Hedgehog signalling pathway or an antagonist of a component of the target pathway of Hedgehog signalling.

- 5 In a further aspect the present invention provides a transgenic animal capable of expressing antagonist of a component of a Hedgehog family member signalling pathway or an antagonist of a component of the target pathway of Hedgehog signalling.
- 10 Various preferred features and embodiments of the present invention will now be described by way of non-limiting example and with reference to the accompanying drawings in which:
- Figure 1 shows a schematic representation of HH signalling;
Figure 2 shows a schematic representation of a component of HH signalling;
- 15 Figure 3 shows a schematic representation of Wnt signalling;
Figure 4 shows plasmid SPC-SV40;
Figure 5 shows plasmid SPC-Shh;
Figure 6 show the results of Example 2;
Figures 7-9 show the results of Example 3;
- 20 Figure 10 show the results of Example 4;
Figure 11 show the results of Example 5;
Figure 12 show the results of Example 6;
Figure 13 show the results of Example 7;
Figure 14 show the results of Example 8;
- 25 Figure 15 show the results of Example 9;
Figure 16 show the results of Example 12; and
Figure 17 show the results of Example 13.

For ease of reference a summary of the accompanying sequence listings is given
30 below:

- SEQ ID NO:1 shows the deduced amino acid sequence of mouse SHH and SEQ ID NO:2 shows the corresponding nucleic acid sequence;
- SEQ ID NO:3 shows the deduced amino acid sequence of mouse Dvl-1 and SEQ ID NO:4 shows the corresponding nucleic acid sequence;
- 5 SEQ ID NO:5 shows the deduced amino acid sequence of mouse HIP and SEQ ID NO:6 shows the corresponding nucleic acid sequence; and
- SEQ ID NO:7 shows the deduced amino acid sequence of mouse WIF-1 and SEQ ID NO:8 shows the corresponding nucleic acid sequence.
- 10 References and Accession Nos. are herein incorporated by reference.

Hedgehog Family Proteins

- All multicellular organisms require cell communication to regulate growth and differentiation in the embryo. One strategy for this is to establish discrete organising centres that emit signals to coordinately control cell proliferation and cell fate determination. The *hedgehog* (*hh*) gene was identified originally through the segment polarity phenotype caused by its mutation in *Drosophila*. Genes of the *hh* family have now been isolated from several vertebrate species,
- 15 including mouse, chicken, zebrafish, rat, *Xenopus* and human. The genes not only seem to show a high degree of structural homology both within and between species, but in addition exhibit some remarkable similarities in the ways in which they function in various embryonic processes. In vertebrates, *Sonic hedgehog* (*Shh*) is a key signal in several signalling centres. There are two other
- 20 mammalian HH members, *Indian hedgehog* (*Ihh*) and *Desert hedgehog* (*Dhh*).

A summary of various *hedgehog* genes is given in the following Table 1:

Table 1

Gene	Species
<i>hedgehog</i> (<i>hh</i>)	<i>Drosophila</i>

<i>Sonic hedgehog (Shh)</i>	Mouse, Human, Rat, <i>Xenopus</i> , Chicken, Zebrafish
<i>Indian hedgehog (Ihh)</i>	Mouse, Human, Chicken
<i>Desert hedgehog (Dhh)</i>	Mouse
<i>Banded hedgehog (X-bhh)</i>	<i>Xenopus</i>
<i>Cepalic hedgehog (X-chh)</i>	<i>Xenopus</i>
<i>tiggy-winkle hedgehog (twhh)</i>	Zebrafish
<i>echidna hedgehog (ehh)</i>	Zebrafish

The classification of genes from different species is based on the comparison of the expression pattern and the amino acid sequence. Of all vertebrate proteins, DHH is most similar to *Drosophila* HH (51% identity over entire length of processed proteins). Amino acid identity among SHH is 93% between mouse and human, 84% between mouse and chicken, 78% between mouse and *Xenopus*, and 68% between mouse and zebrafish. Intraspecies comparison within the mouse reveals 58-63% identity in pairwise combination between SHH, IHH and DHH. Interspecies comparison between the mouse and *Xenopus* reveals highest identities between IHH and XBHH (70%) and DHH and XCHH (64%).

The various Hedgehog proteins consist of a signal peptide, with a highly conserved N-terminal region and a more divergent C-terminal domain. It is understood that the biologically active Hedgehog peptides are formed from a larger precursor protein. In addition to signal sequence cleavage in the secretory pathway, Hedgehog precursor proteins undergo an internal autoproteolytic cleavage. This autocleavage generates an N-terminal peptide (about 19kDa) and a C-terminal peptide (of about 26-28kDa). It is this N-terminal peptide that is necessary for short- and long-range Hedgehog signalling activities in *Drosophila* and vertebrates. The N-terminal peptide stays tightly associated with the surface of cells in which it is synthesised, while the C-terminal peptide is freely diffusible.

Signalling Pathway

Figure 1 shows one representation of a Hedgehog signalling pathway, with
5 particular reference to signalling in vertebrates.

Epithelial cells may express the homeodomain transcription factor engrailed (*en*)
and secrete Hedgehog protein shown for illustrative purposes in the Figure as
Shh. We have observed that En plays an important role in the maintainance of
10 lymphocyte survival in the peripheral immune system.

In target cells, HH signalling is mediated by two transmembrane proteins
patched (Ptc) which has structural similarities to channel and transporter
proteins, and Smoothened (Smo), a seven-transmembrane protein similar to G-
15 protein coupled receptors and the Wingless receptor Frizzled (described below).
Smo is a constitutive activator of HH target genes. Its activity is normally
repressed by Ptc, and this repression is relieved by HH binding to Ptc. Thus,
binding of HH to Ptc allows signal transduction leading to activation of the
transcription factor Gli, which is located in the nucleus of the target cells.

20 The signal reaches Gli through the cytoplasmic complex formed between (1) the
serine/threonine kinase Fused (Fu), (2) Suppressor of Fused (SU(Fu)); and (3)
Costal2 (Cos2). Signalling through this complex may be inhibited by the cAMP-
dependent protein kinase A (PKA) (see Figure 2).

25 Gli acts on target genes wingless (Wnt) and the BMP /activin growth factors.
Both Wnt and BMP are secreted to the extracellular fluid to bind to their
receptors. This process is illustrated schematically in Figure 1.

A summary and comparison of components of the Hedgehog signalling pathway is given below in Table 2:

Table 2

<i>Drosophila</i>	Vertebrate
En	En 1,2
hh	Ihh, Dhh, Shh
Ptc	Ptc 1,2
Smo	Smo
Ci	Gli 1-3
Target genes	
Wg	Wnt ~15
Dpp \equiv TGF β	BMP 8-10

5

The nomenclature may be used interchangeably herein.

Further information on Hedgehog signalling may be found in the following articles: Ingham; Chuang and McMahon; Picicelli *et al*; and Hammerschmidt *et al*.

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BMP signalling pathway

Bone morphogenic proteins (BMPs) are multifunctional cytokines, which are members of the transforming growth factor- β (TGF- β) superfamily. They regulate cellular proliferation, differentiation, apoptosis of various cells types. Activities of BMPs are extracellularly regulated by BMP-binding proteins, Noggin, Chordin, Gremlin Cerberus and Xnr-3. BMPs have been found to block neurogenesis in early development, but this can be resolved by soluble factors, e.g. Noggin and Chordin, which are secreted extracellularly and which bind and

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neutralize the BMPs preventing them from activating their receptors. In addition, signalling through BMP receptors can inhibit expression of the Notch ligand Delta, which we have previously shown to be important in regulating peripheral immune responses. The Notch signalling pathway is discussed in our
5 International patent publication No. WO98/20142.

BMPs bind to two different types of serine-threonine kinase receptors, type I and type II. As for type I receptors, BMPs bind to BMP type IA receptor, BMP type IB receptor and activin type I receptor. As for type II receptors, BMPs bind to
10 BMP type II receptor, activin type II receptor and activin type IIB receptor. In the receptor-ligand complexes, type II receptors phosphorylate type I receptors in the GS domain (rich in glycine, serine and threonine residues) to activate the latter. The activated type I receptors phosphorylate Smad family, which transduces the signals from cytoplasm into nuclei. Smad1, Smad5 and possibly
15 MADH6 are activated by BMP receptors, form heteromeric complexes with Smad4, and translocate into the nucleus where they may activate transcription of various genes. Smad6 and Smad7 are inhibitory Smads, and may act as autocrine switch-off signals. BMP induced Smad signalling down regulates achaete/scute gene expression which is required for expression of the Notch ligand Delta. As
20 indicated in Table 2 above, in *Drosophila*, Decapentaplegic (Dpp) is a homolog of mammalian BMPs.

Wingless/Wnt signalling pathway

25 We have examined the role for dysregulation of the Wnt signalling pathway in interstitial lung disease. The Wnt genes are targets of the HH pathway, and the Wnt proteins are secreted growth factors which are involved in the regulation of epithelial cell proliferation and differentiation in the lung during embryonic development. We propose that Wnt signalling may also be upregulated during
30 processes of epithelial cell repair in the lung.

Dishevelled-1 (Dvl-1) is the murine homolog of the fly Dsh gene and functions to transmit signals from the Wnt receptor, Frizzled, to the cytoplasm, where it regulates the kinase activity of a well known serine/threonine kinase, GSK-3b.

5 Over expression of Dsh in fly epithelia leads to oncogenic activation of the epithelium by increasing Wnt signalling.

A representation of this pathway is shown in Figure 3. Wingless (Wg), in *Drosophila*, and, its vertebrate homolog, Wnt signalling pathways regulate cell

10 proliferation. Wg and Wnt are secreted growth factors which are involved in triggering cellular decisions. The Wg/Wnt ligand binds to Frizzled (Fz) family receptor molecules to initiate a signal transduction cascade involving the cytoplasmic protein Dishevelled (Dvl) (Sussman DJ *et al*). The GenBank accession number for *Dvl-1* cDNA is U10115. The complex illustrated in Figure

15 3 is present in the cytoplasm of the target cell. Generally APC blocks signalling; however, in the presence of signalling from Wnt, β -catenin is released and interacts with two transcription factors - Lef-1/TCF-1 resulting in target gene expression. Target genes of Wnt include En and therefore indirectly HH, c-myc and cyclin D1. It will be appreciated that Notch signalling is also regulated by

20 the Wnt pathway, as Dvl has been found to inhibit Notch signalling.

Inhibitors

The present invention relates to the use of compounds which inhibit or block

25 (antagonise) Hedgehog signalling. Such compounds may be seen as having the effect of downregulating the expression of Hedgehog. Similarly the present invention also relates to the use of compounds which inhibit or block (antagonise) a signalling pathway which is a target of the Hedgehog signalling pathway. Conveniently such compounds may be referred to herein as inhibitors

30 or antagonists.

The invention contemplates that mutations that result in loss of normal function of the regulators of the Hedgehog signalling pathway or regulators of a pathway which is a target of the Hedgehog signalling pathway in human disease states in which lymphocyte infiltration or failure of a cell cycle checkpoint is involved. Gene therapy to restore such regulatory activity would thus be indicated in treating those disease states. Alternatively, it is contemplated that preventing the expression of or inhibiting the activity of such signalling pathways will be useful in treating the disease states. It is contemplated that antisense therapy or gene therapy could be applied to negatively regulate such signalling pathways.

Antagonists for each component of the signalling pathway have been identified. These may be summarised as follows in Table 3:

Table 3

Component	Antagonist
HH	Hip (Chuang and McMahon), Veratrum alkaloids and distal inhibitors of cholesterol biosynthesis (Cooper <i>et al</i>) e.g. cyclopamine (Coventry <i>et al</i>).
Wnt	Fzb (Leyns <i>et al</i>), Cerberus (Bouwmeester <i>et al</i>), Gremlin (Hsu <i>et al</i>), WIF-1 (Hsieh <i>et al</i>)
BMP	Noggin (Valenzuela <i>et al</i>), Chordin (Sasai <i>et al</i>), Cerberus, Gremlin, Xnr-3
Activin	Follistatin (Iemura <i>et al</i>)

HIP (for Hedgehog-interacting protein) is a membrane glycoprotein that binds to at least all three mammalian Hedgehog proteins with an affinity comparable to that of Ptc. HIP appears to attenuate Hedgehog signalling as a result of binding to Hedgehog proteins. Such a negative regulatory feedback loop could also serve

to modulate the response to any Hedgehog signal. The GenBank accession number for HIP is AF116865.

Veratrum alkaloids and distal inhibitors of cholesterol biosynthesis have been studied for more than 30 years as potent teratogens capable of inducing cyclopia and other birth defects. It has also been shown that these compounds specifically block the Shh signaling pathway (Cooper *et al*). One example of such a veratrum alkaloid is cyclopamine (11-deoxojervine), a steroid isolated from the desert plant *Veratrum californicum* (Coventry *et al*).

10

Fzb (Frezzed) is a secreted antagonist of Wnt signalling. Fzb contains a domain similar to the putative Wnt-binding region of the Frizzled family of transmembrane receptors, but it lacks all the transmembrane domains resulting in a putative secreted Wnt-binding protein. The GenBank accession numbers for the *Xenopus*, mouse and human Fzb cDNA sequences are U68059, U68058 and U68057, respectively.

15

Cerberus is a secreted protein and it has been found to be an antagonist of the Wnt and BMP signalling pathways. The GenBank accession number for the *Xenopus* Cerberus cDNA is U64831.

20

WIF-1 (Wnt-inhibitory factor-1) is a secreted protein which binds to Wnt proteins and inhibitors their activities. GenBank accession numbers for WIF-1 are: human, AF122922; mouse, AF122923; *Xenopus*, AF122924; and zebrafish, AF122925.

25

Noggin and Chordin bind to BMPs thereby preventing activation of their signalling cascade.

Gremlin is a secreted protein and it has been found to be an antagonist of the Wnt and BMP signalling pathways. The GenBank accession numbers for Gremlin cDNA are: *Xenopus*, AF045798; chick, AF045799; human, AF045800; and mouse, AF045801.

5

Xnr-3 has been found to be an antagonist of BMP signalling pathways.

Follistatin has been found to inhibit others aspects of BMP activity as well as acting as an activin-binding protein.

10

It will also be appreciated that the antagonist may itself be a component of the Hedgehog signalling pathway, or a component of the target pathway of the Hedgehog signalling pathway. Examples of such antagonists include the negative regulators of HH signalling: Ptc, Cos2 and PKA.

15

In a particularly preferred embodiment use is made of PKA. PKA has been implicated in the mechanism of Hh signal transduction because it acts to repress Hh target genes in imaginal disc cells that express Ci. Ci action as transcriptional repressor or activator is contingent upon Hedgehog-regulated,

20

PKA-dependent proteolytic processing.

Cyclic AMP (cAMP) is a nucleotide that is generated from ATP in response to hormonal stimulation of cell-surface receptors. cAMP acts as a signaling molecule by activating A-kinase; it is hydrolyzed to AMP by phosphodiesterase (PDE). cAMP levels affect cubitus cleavage and TGF- β levels. Specifically, when cAMP levels increase, TGF- β levels decrease and this will affect fibrosis, for example. In another embodiment of the invention use is made of cAMP modifiers in treatment. Such modifiers include PDE inhibitors, and beta-agonists such as the beta-adrenergic agonist. For example, it has been found that

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ptc 1 transcription can be induced by agents activating the cAMP signal transduction pathway.

Antisense nucleic acids (preferably 10 to 20 base pair oligonucleotides) capable of specifically binding to expression control sequences or RNA are introduced into cells (*e.g.*, by a viral vector or colloidal dispersion system such as a liposome). The antisense nucleic acid binds to the target sequence in the cell and prevents transcription or translation of the target sequence. Phosphothioate and methylphosphate antisense oligonucleotides are specifically contemplated for therapeutic use by the invention. The antisense oligonucleotides may be further modified by poly-L-lysine, transferrin polylysine, or cholesterol moieties at their 5' end.

Also comprehended by the present invention are antibody products (*e.g.*, monoclonal and polyclonal antibodies, single chain antibodies, chimeric antibodies, CDR-grafted antibodies and antigen-binding fragments thereof) and other binding proteins (such as those identified in the assays above). Binding proteins can be developed using isolated natural or recombinant enzymes. The binding proteins are useful, in turn, for purifying recombinant and naturally occurring enzymes and identifying cells producing such enzymes. Assays for the detection and quantification of proteins in cells and in fluids may involve a single antibody substance or multiple antibody substances in a "sandwich" assay format to determine cytological analysis of HH protein levels. The binding proteins are also manifestly useful in modulating (*i.e.* blocking, inhibiting, or stimulating) interactions.

Antibodies may be generated by administering polypeptides or epitope-containing fragments to an animal, usually a rabbit, using routine protocols. Examples of such techniques include those in Kohler and Milstein.

More generally, the antagonist may be selected from polypeptides and fragments thereof, linear peptides, cyclic peptides, synthetic and natural compounds including low molecular weight organic or inorganic compounds. The antagonist may be derived from a biological material such as a component of extracellular matrix.

Polypeptide substances, such as Noggin or Chordin, may be purified from mammalian cells, obtained by recombinant expression in suitable host cells or obtained commercially. Alternatively, nucleic acid constructs encoding the polypeptides may be introduced by transfection using standard techniques or viral infection/transduction.

Inhibitors for use according to the present invention may be conveniently identified using a convenient screening procedure.

15

One assay for identifying such inhibitors may involve immobilizing a component of the relevant pathway, e.g. HH, or a test protein, detectably labelling the nonimmobilized binding partner, incubating the binding partners together and determining the amount of label bound. Bound label indicates that the test protein interacts with the component.

20

Another type of assay for identifying inhibitors involves immobilizing a component of the pathway, e.g. HH, or a fragment thereof on a solid support coated (or impregnated with) a fluorescent agent, labelling a test protein with a compound capable of exciting the fluorescent agent, contacting the immobilized component with the labelled test protein, detecting light emission by the fluorescent agent, and identifying interacting proteins as test proteins which result in the emission of light by the fluorescent agent. Alternatively, the putative interacting protein may be immobilized and the component may be labelled in the assay.

25
30

Moreover, such assays for identifying inhibitors may involve: transforming or transfecting appropriate host cells with a DNA construct comprising a reporter gene under the control of a promoter regulated by a transcription factor having a DNA-binding domain and an activating domain; expressing in the host cells a first hybrid DNA sequence encoding a first fusion of part or all of a component of the pathway, e.g. HH, Wnt or BMP, and the DNA binding domain or the activating domain of the transcription factor; expressing in the host cells a second hybrid DNA sequence encoding part or all of a protein that interacts with said component and the DNA binding domain or activating domain of the transcription factor which is not incorporated in the first fusion; evaluating the effect of a test compound on the interaction between said component and the interacting protein by detecting binding of the interacting protein to said component in a particular host cell by measuring the production of reporter gene product in the host cell in the presence or absence of the test compound; and identifying modulating compounds as those test compounds altering production of the reported gene product in comparison to production of the reporter gene product in the absence of the modulating compound. Presently preferred for use in the assay are a *lexA* promoter to drive expression of the reporter gene, the *lacZ* reporter gene, a transcription factor comprising the *lexA* DNA binding domain and the GAL4 transactivation domain, and yeast host cells.

In a particular embodiment described in relation to Hedgehog signalling the appropriate host cell is transformed or transfected with a DNA construct comprising a reporter gene under the control of the Ptc promoter; expressing in said cells a DNA sequence encoding Hedgehog; evaluating the effect of a test compound on the interaction between HH and the Ptc promoter in a particular host cell by measuring the production of reporter gene product in the host cell in the absence and presence of the test compound; and identifying inhibitors as those test compounds reducing the production of the reporter gene product in

comparison to production of the reporter gene product in the absence of the test compound.

5 Analogous assays may be used for inhibitors of the target pathways of Hedgehog signalling. For example, for the Wnt signalling pathway, the ability of a compound to modulate the interaction of Wnt and Fz may be determined. For BMP signalling the ability of a compound to modulate the interaction of BMP and its BMP receptor may be determined.

10 Combinatorial libraries, peptide and peptide mimetics, defined chemical entities, oligonucleotides, and natural product libraries may be screened for activity as inhibitors in such assays.

15 The present invention also relates to the use of derivatives, variants, fragments, analogues, homologues and mimetics of the inhibitors mentioned above, including those identifiable using the assay procedures.

20 The term "derivative" as used herein in relation to the polypeptides of the present invention includes any substitution of, variation of, modification of, replacement of, deletion of, or addition of one (or more) amino acid residues from or to the sequence providing that the resultant protein etc., possesses the capability to antagonise the action of the signalling pathway.

25 The term "variant" as used herein in relation to the polypeptides of the present invention includes any substitution of, variation of, modification of, replacement of, deletion of, or addition of one (or more) amino acid residues from or to the sequence providing that the resultant protein etc., possesses the capability to antagonise the action of the signalling pathway.

The term "fragment" as used herein in relation to the polypeptides of the present invention includes a variant polypeptide which has an amino acid sequence that is entirely the same as part but not all of the amino acid sequence of the aforementioned polypeptide and possesses the capability to antagonise the action
5 of the signalling pathway.

The term "analogue" as used herein in relation to the polypeptides of the present invention includes any peptidomimetic, i.e. a chemical compound that possess the capability to antagonise the action of the signalling pathway in a similar
10 manner to the parent polypeptide.

The term "homologue" as used herein in relation to the polypeptides of the present invention includes a polypeptide which has the same evolutionary origin as the subject polypeptide providing that it possesses the capability to antagonise
15 the action of the signalling pathway.

The term "mimetic" as used herein in relation to the inhibitors of the present invention includes a compound which also possesses the capability to antagonise the action of the signalling pathway in a similar manner to the parent compound.
20

More particularly, the term "homologue" covers identity with respect to structure and/or function providing the expression product of the resultant nucleotide sequence has the inhibitory activity. With respect to sequence identity (i.e. similarity), preferably there is at least 75%, more preferably at least 85%, more
25 preferably at least 90% sequence identity. More preferably there is at least 95%, more preferably at least 98%, sequence identity. These terms also encompass allelic variations of the sequences.

Sequence identity with respect to the sequences can be determined by a simple
30 "eyeball" comparison (i.e. a strict comparison) of any one or more of the

sequences with another sequence to see if that other sequence has, for example, at least 75% sequence identity to the sequence(s).

Relative sequence identity can also be determined by commercially available
5 computer programs that can calculate % identity between two or more sequences using any suitable algorithm for determining identity, using for example default parameters. A typical example of such a computer program is CLUSTAL. Advantageously, the BLAST algorithm is employed, with parameters set to default values. The BLAST algorithm is described in detail at
10 http://www.ncbi.nih.gov/BLAST/blast_help.html, which is incorporated herein by reference. The search parameters are defined as follows, can be advantageously set to the defined default parameters.

Advantageously, "substantial identity" when assessed by BLAST equates to
15 sequences which match with an EXPECT value of at least about 7, preferably at least about 9 and most preferably 10 or more. The default threshold for EXPECT in BLAST searching is usually 10.

BLAST (Basic Local Alignment Search Tool) is the heuristic search algorithm
20 employed by the programs blastp, blastn, blastx, tblastn, and tblastx; these programs ascribe significance to their findings using the statistical methods of Karlin and Altschul (see http://www.ncbi.nih.gov/BLAST/blast_help.html) with a few enhancements. The BLAST programs were tailored for sequence similarity searching, for example to identify homologues to a query sequence.
25 For a discussion of basic issues in similarity searching of sequence databases, see Altschul *et al* (1994) Nature Genetics 6:119-129.

The five BLAST programs available at <http://www.ncbi.nlm.nih.gov> perform the following tasks:

30

blastp - compares an amino acid query sequence against a protein sequence database.

5 **blastn** - compares a nucleotide query sequence against a nucleotide sequence database.

blastx - compares the six-frame conceptual translation products of a nucleotide query sequence (both strands) against a protein sequence database.

10 **tblastn** - compares a protein query sequence against a nucleotide sequence database dynamically translated in all six reading frames (both strands).

tblastx - compares the six-frame translations of a nucleotide query sequence against the six-frame translations of a nucleotide sequence database.

15 BLAST uses the following search parameters:

HISTOGRAM - Display a histogram of scores for each search; default is yes.
(See parameter H in the BLAST Manual).

20 DESCRIPTIONS - Restricts the number of short descriptions of matching sequences reported to the number specified; default limit is 100 descriptions.
(See parameter V in the manual page).

EXPECT - The statistical significance threshold for reporting matches against
25 database sequences; the default value is 10, such that 10 matches are expected to be found merely by chance, according to the stochastic model of Karlin and Altschul (1990). If the statistical significance ascribed to a match is greater than the EXPECT threshold, the match will not be reported. Lower EXPECT thresholds are more stringent, leading to fewer chance matches being reported.

30 Fractional values are acceptable. (See parameter E in the BLAST Manual).

- CUTOFF - Cutoff score for reporting high-scoring segment pairs. The default value is calculated from the EXPECT value (see above). HSPs are reported for a database sequence only if the statistical significance ascribed to them is at least
5 as high as would be ascribed to a lone HSP having a score equal to the CUTOFF value. Higher CUTOFF values are more stringent, leading to fewer chance matches being reported. (See parameter S in the BLAST Manual). Typically, significance thresholds can be more intuitively managed using EXPECT.
- 10 ALIGNMENTS - Restricts database sequences to the number specified for which high-scoring segment pairs (HSPs) are reported; the default limit is 50. If more database sequences than this happen to satisfy the statistical significance threshold for reporting (see EXPECT and CUTOFF below), only the matches ascribed the greatest statistical significance are reported. (See parameter B in the
15 BLAST Manual).
- MATRIX - Specify an alternate scoring matrix for BLASTP, BLASTX, TBLASTN and TBLASTX. The default matrix is BLOSUM62 (Henikoff & Henikoff, 1992). The valid alternative choices include: PAM40, PAM120,
20 PAM250 and IDENTITY. No alternate scoring matrices are available for BLASTN; specifying the MATRIX directive in BLASTN requests returns an error response.
- STRAND - Restrict a TBLASTN search to just the top or bottom strand of the
25 database sequences; or restrict a BLASTN, BLASTX or TBLASTX search to just reading frames on the top or bottom strand of the query sequence.
- FILTER - Mask off segments of the query sequence that have low compositional complexity, as determined by the SEG program of Wootton & Federhen (1993)
30 Computers and Chemistry 17:149-163, or segments consisting of short-

periodicity internal repeats, as determined by the XNU program of Claverie & States (1993) Computers and Chemistry 17:191-201, or, for BLASTN, by the DUST program of Tatusov and Lipman (see <http://www.ncbi.nlm.nih.gov>). Filtering can eliminate statistically significant but biologically uninteresting reports from the blast output (e.g., hits against common acidic-, basic- or proline-rich regions), leaving the more biologically interesting regions of the query sequence available for specific matching against database sequences.

Low complexity sequence found by a filter program is substituted using the letter "N" in nucleotide sequence (e.g., "NNNNNNNNNNNNNN") and the letter "X" in protein sequences (e.g., "XXXXXXXXXX").

Filtering is only applied to the query sequence (or its translation products), not to database sequences. Default filtering is DUST for BLASTN, SEG for other programs.

It is not unusual for nothing at all to be masked by SEG, XNU, or both, when applied to sequences in SWISS-PROT, so filtering should not be expected to always yield an effect. Furthermore, in some cases, sequences are masked in their entirety, indicating that the statistical significance of any matches reported against the unfiltered query sequence should be suspect.

NCBI-gi - Causes NCBI gi identifiers to be shown in the output, in addition to the accession and/or locus name.

Most preferably, sequence comparisons are conducted using the simple BLAST search algorithm provided at <http://www.ncbi.nlm.nih.gov/BLAST>.

Other computer program methods to determine identify and similarity between the two sequences include but are not limited to the GCG program package

(Devereux *et al* 1984 Nucleic Acids Research 12: 387) and FASTA (Atschul *et al* 1990 J Molec Biol 403-410).

5 In some aspects of the present invention, no gap penalties are used when determining sequence identity.

The present invention also encompasses use of nucleotide sequences that are complementary to the sequences presented herein, or any fragment or derivative thereof. If the sequence is complementary to a fragment thereof then that sequence
10 can be used as a probe to identify similar promoter sequences in other organisms.

The present invention also encompasses use of nucleotide sequences that are capable of hybridising to the sequences presented herein, or any fragment or derivative thereof.

15 Hybridization means a "process by which a strand of nucleic acid joins with a complementary strand through base pairing" (Coombs J (1994) Dictionary of Biotechnology, Stockton Press, New York NY) as well as the process of amplification as carried out in polymerase chain reaction technologies as described in Dieffenbach CW and GS Dveksler (1995, PCR Primer, a
20 Laboratory Manual, Cold Spring Harbor Press, Plainview NY).

Also included within the scope of the present invention are use of nucleotide sequences that are capable of hybridizing to the nucleotide sequences presented
25 herein under conditions of intermediate to maximal stringency. Hybridization conditions are based on the melting temperature (T_m) of the nucleic acid binding complex, as taught in Berger and Kimmel (1987, Guide to Molecular Cloning Techniques, Methods in Enzymology, Vol 152, Academic Press, San Diego CA), and confer a defined "stringency" as explained below.

30

Maximum stringency typically occurs at about $T_m - 5^\circ\text{C}$ (5°C below the T_m of the probe); high stringency at about 5°C to 10°C below T_m ; intermediate stringency at about 10°C to 20°C below T_m ; and low stringency at about 20°C to 25°C below T_m . As will be understood by those of skill in the art, a maximum stringency hybridization can be used to identify or detect identical nucleotide sequences while an intermediate (or low) stringency hybridization can be used to identify or detect similar or related nucleotide sequences.

In a preferred aspect, the present invention covers use of nucleotide sequences that can hybridise to the nucleotide sequences of the present invention under stringent conditions (e.g. 65°C and $0.1\times\text{SSC}$).

The present invention also encompasses use of nucleotide sequences that are capable of hybridising to the sequences that are complementary to the sequences presented herein, or any fragment or derivative thereof. Likewise, the present invention encompasses use of nucleotide sequences that are complementary to sequences that are capable of hybridising to the sequence of the present invention. These types of nucleotide sequences are examples of variant nucleotide sequences.

In this respect, the term "variant" encompasses sequences that are complementary to sequences that are capable of hybridising to the nucleotide sequences presented herein. Preferably, however, the term "variant" encompasses sequences that are complementary to sequences that are capable of hybridising under stringent conditions (eg. 65°C and $0.1\times\text{SSC}$ { $1\times\text{SSC} = 0.15\text{ M NaCl}$, $0.015\text{ Na}_3\text{ citrate pH } 7.0$ }) to the nucleotide sequences presented herein.

Transgenic Animals

The present invention also relates to transgenic animals which are capable of expressing or overexpressing at least one antagonist useful in the present

invention. Preferably the animal expresses or overexpresses HIP, Fzb-1, Noggin (Ngg) and/or WIF-1.

5 The present invention additionally relates to transgenic animals which are capable of expressing or overexpressing at least one polypeptide which is a component of the Hedgehog signalling pathway or a component of a pathway which is a target of the Hedgehog signalling pathway, such as the Wnt or BMP signalling pathway. Preferably the animal expresses or overexpresses HH (more preferably Shh), and/or Dvl-1.

10

The transgenic animal is typically a vertebrate, more preferably a rodent, such as a rat or a mouse, but also includes other mammals such as human, goat, pig or cow etc.

15 Such transgenic animals are useful as animal models of disease and in screening assays for new useful compounds. By specifically expressing one or more polypeptides, as defined above, the effect of such polypeptides on the development of disease can be studied. Furthermore, therapies including gene therapy and various drugs can be tested on transgenic animals. Methods for the
20 production of transgenic animals are known in the art. For example, there are several possible routes for the introduction of genes into embryos. These include (i) direct transfection or retroviral infection of embryonic stem cells followed by introduction of these cells into an embryo at the blastocyst stage of development; (ii) retroviral infection of early embryos; and (iii) direct microinjection of DNA
25 into zygotes or early embryo cells.

The present invention also includes stable cell lines for use as disease models for testing or treatment.

A stable cell line will contain a recombinant gene or genes, also known herein as a transgene, encoding one or more inhibitors or components of a Hedgehog signalling pathway or of a pathway which is a target of the Hedgehog signalling pathway.

5

Preferably the transgene is HH (more preferably Shh), HIP, WIF-1, Fzb-1, Ngg and/or Dvl-1. A cell line containing a transgene, as described herein, is made by introducing the transgene into a selected cell line according to one of several procedures known in the art for introducing a foreign gene into a cell.

10

As also described below, the sequences encoding the inhibitors and components of signalling pathways, as described herein, are operably linked to control sequences, including promoters/enhancers and other expression regulation signals.

15

The promoter is typically selected from promoters which are functional in mammalian cells, although prokaryotic promoters and promoters functional in other eukaryotic cells may be used. The promoter is typically derived from promoter sequences of viral or eukaryotic genes. For example, it may be a promoter derived from the genome of a cell in which expression is to occur. With respect to eukaryotic promoters, they may be promoters that function in a ubiquitous manner (such as promoters of α -actin, β -actin, tubulin) or, alternatively, a tissue-specific manner (such as promoters of the genes for pyruvate kinase). Tissue-specific promoters specific for lymphocytes, dendritic cells, skin, brain cells and epithelial cells within the eye are particularly preferred, for example the CD2, CD11c, keratin 14, Wnt-1 and Rhodopsin promoters respectively. Preferably the epithelial cell promoter SPC is used. They may also be promoters that respond to specific stimuli, for example promoters that bind steroid hormone receptors. Viral promoters may also be used, for example the Moloney murine leukaemia virus long terminal repeat

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30

(MMLV LTR) promoter, the rous sarcoma virus (RSV) LTR promoter or the human cytomegalovirus (CMV) IE promoter.

5 It may also be advantageous for the promoters to be inducible so that the levels of expression of the heterologous gene can be regulated during the life-time of the cell. Inducible means that the levels of expression obtained using the promoter can be regulated.

10 In addition, any of these promoters may be modified by the addition of further regulatory sequences, for example enhancer sequences. Chimeric promoters may also be used comprising sequence elements from two or more different promoters described above.

Therapeutic Uses

15

As previously mentioned, in many tissues, such as lung and kidney, inflammation may lead to chronic diseases. For example, chronic lung disease involve tissue remodelling in the presence of inflammatory cells, examples are emphysema and interstitial lung disease (ILD). The disease tissue is associated with epithelial cell hyperplasia leading to fibrosis and scarring. There is an accompanying mononuclear cell infiltration at local sites of inflammation and the induction of immune responses reactive with self antigens. Similar pathology is also observed in the chronic rejection of graft tissue, including transplanted organs.

25

Hedgehog is important in regulating growth and differentiation of epithelial cells. It has a role in the formation of notochord, limb, gut, lung, skin etc. Branching morphogenesis occurs through induction of Wnt and BMP growth factors. It binds to its receptor Ptc and Smo. Mutations can lead to the human autosomal disorder Nevoid basal cell carcinoms syndrome (NBCCS) which is

30

characterised by developmental abnormalities and a high predisposition for various forms of cancer mainly the very common basal cell carcinomas (BCC).

5 BMPs are members of the TGF- β superfamily. Whereas a role for TGF- β 1 in mediating lung fibrosis is well established, previously BMP-4 has only been implicated in fibrotic disease of bone.

We now provide a method for the treatment of epithelial cell hyperplasia and fibrosis particularly in the lung and kidney. More particularly diseases which
10 may be treated include adult respiratory distress syndrome; chronic obstructive airway disorders/ chronic obstructive pulmonary disease including asthma, emphysema and chronic bronchitis; atelectasis; occupational lung disease including silicosis; hypersensitivity diseases of the lung including hypersensitivity pneumonitis; idiopathic interstitial lung diseases including
15 idiopathic pulmonary fibrosis, usual interstitial pneumonia, desquamative interstitial pneumonia and acute interstitial pneumonia; and pleural fibrosis. Further details on such conditions and those given below may be found in The Merck Manual (17th Edition), published by Merck Research Laboratories, N.J., USA.

20

The present invention is also useful in treating immune disorders such as autoimmune diseases or graft rejection such as allograft rejection.

Examples of disorders that may be treated include a group commonly called
25 autoimmune diseases. The spectrum of autoimmune disorders ranges from organ specific diseases (such as thyroiditis, insulinitis, multiple sclerosis, iridocyclitis, uveitis, orchitis, hepatitis, Addison's disease, myasthenia gravis) to systemic illnesses such as rheumatoid arthritis or lupus erythematosus. Other disorders include immune hyperreactivity, such as allergic reactions.

30

In more detail: Organ-specific autoimmune diseases include multiple sclerosis, insulin dependent diabetes mellitus, several forms of anemia (aplastic, hemolytic), autoimmune hepatitis, thyroiditis, insulinitis, iridocyclitis, skleritis, uveitis, orchitis, myasthenia gravis, idiopathic thrombocytopenic purpura,
5 inflammatory bowel diseases (Crohn's disease, ulcerative colitis).

Systemic autoimmune diseases include: rheumatoid arthritis, juvenile arthritis, scleroderma and systemic sclerosis, sjogren's syndrom, undifferentiated connective tissue syndrome, antiphospholipid syndrome, different forms of
10 vasculitis (polyarteritis nodosa, allergic granulomatosis and angiitis, Wegner's granulomatosis, Kawasaki disease, hypersensitivity vasculitis, Henoch-Schoenlein purpura, Behcet's Syndrome, Takayasu arteritis, Giant cell arteritis, Thrombangiitis obliterans), lupus erythematosus, polymyalgia rheumatica, essentiell (mixed) cryoglobulinemia, Psoriasis vulgaris and psoriatic arthritis,
15 diffus fasciitis with or without eosinophilia, polymyositis and other idiopathic inflammatory myopathies, relapsing panniculitis, relapsing polychondritis, lymphomatoid granulomatosis, erythema nodosum, ankylosing spondylitis. Reiter's syndrome, different forms of inflammatory dermatitis.

20 A more extensive list of disorders includes: unwanted immune reactions and inflammation including arthritis, including rheumatoid arthritis, inflammation associated with hypersensitivity, allergic reactions, asthma, systemic lupus erythematosus, collagen diseases and other autoimmune diseases, inflammation associated with atherosclerosis, arteriosclerosis, atherosclerotic heart disease,
25 reperfusion injury, cardiac arrest, myocardial infarction, vascular inflammatory disorders, respiratory distress syndrome or other cardiopulmonary diseases, inflammation associated with peptic ulcer, ulcerative colitis and other diseases of the gastrointestinal tract, hepatic fibrosis, liver cirrhosis or other hepatic diseases, thyroiditis or other glandular diseases, glomerulonephritis or other
30 renal and urologic diseases, otitis or other oto-rhino-laryngological diseases,

dermatitis or other dermal diseases, periodontal diseases or other dental diseases, orchitis or epididymo-orchitis, infertility, orchidal trauma or other immune-related testicular diseases, placental dysfunction, placental insufficiency, habitual abortion, eclampsia, pre-eclampsia and other immune and/or

5 inflammatory-related gynaecological diseases, posterior uveitis, intermediate uveitis, anterior uveitis, conjunctivitis, chorioretinitis, uveoretinitis, optic neuritis, intraocular inflammation, e.g. retinitis or cystoid macular oedema, sympathetic ophthalmia, scleritis, retinitis pigmentosa, immune and inflammatory components of degenerative fundus disease, inflammatory

10 components of ocular trauma, ocular inflammation caused by infection, proliferative vitreo-retinopathies, acute ischaemic optic neuropathy, excessive scarring, e.g. following glaucoma filtration operation, immune and/or inflammation reaction against ocular implants and other immune and inflammatory-related ophthalmic diseases, inflammation associated with

15 autoimmune diseases or conditions or disorders where, both in the central nervous system (CNS) or in any other organ, immune and/or inflammation suppression would be beneficial, Parkinson's disease, complication and/or side effects from treatment of Parkinson's disease, AIDS-related dementia complex HIV-related encephalopathy, Devic's disease, Sydenham chorea, Alzheimer's

20 disease and other degenerative diseases, conditions or disorders of the CNS, inflammatory components of stokes, post-polio syndrome, immune and inflammatory components of psychiatric disorders, myelitis, encephalitis, subacute sclerosing pan-encephalitis, encephalomyelitis, acute neuropathy, subacute neuropathy, chronic neuropathy, Guillain-Barre syndrome, Sydenham

25 chora, myasthenia gravis, pseudo-tumour cerebri, Down's Syndrome, Huntington's disease, amyotrophic lateral sclerosis, inflammatory components of CNS compression or CNS trauma or infections of the CNS, inflammatory components of muscular atrophies and dystrophies, and immune and inflammatory related diseases, conditions or disorders of the central and

30 peripheral nervous systems, post-traumatic inflammation, septic shock,

infectious diseases, inflammatory complications or side effects of surgery or organ, inflammatory and/or immune complications and side effects of gene therapy, e.g. due to infection with a viral carrier, or inflammation associated with AIDS, to suppress or inhibit a humoral and/or cellular immune response, to treat
5 or ameliorate monocyte or leukocyte proliferative diseases, e.g. leukaemia, by reducing the amount of monocytes or lymphocytes, for the prevention and/or treatment of graft rejection in cases of transplantation of natural or artificial cells, tissue and organs such as cornea, bone marrow, organs, lenses, pacemakers, natural or artificial skin tissue.

10

The present invention is also useful in cancer therapy, particularly in diseases involving the conversion of epithelial cells to cancer. The present invention is especially useful in relation to adenocarcinomas such as: small cell lung cancer, and cancer of the kidney, uterus, prostate, bladder, ovary, colon and breast.

15

We have now found that the use of antagonists of Hedgehog signalling may prevent and/or promote regression of the above-mentioned diseases.

Vectors, host cells, expression

20

The present invention also relates to vectors which comprise a polynucleotide useful in the present invention, host cells which are genetically engineered with vectors of the invention and the production of polypeptides useful in the present invention by such techniques.

25

For recombinant production, host cells can be genetically engineered to incorporate expression systems or polynucleotides of the invention. Introduction of a polynucleotide into the host cell can be effected by methods described in many standard laboratory manuals, such as Davis *et al* and Sambrook *et al*, such
30 as calcium phosphate transfection, DEAE-dextran mediated transfection,

transfection, microinjection, cationic lipid-mediated transfection, electroporation, transduction, scrape loading, ballistic introduction and infection.

Representative examples of appropriate hosts include bacterial cells, such as
5 streptococci, staphylococci, *E. Coli*, streptomyces and *Bacillus subtilis* cells;
fungal cells, such as yeast cells and *Aspergillus* cells; insect cells such as
Drosophila S2 and *Spodoptera* Sf9 cells; animal cells such as CHO, COS, HeLa,
C127, 3T3, BHK, 293 and Bowes melanoma cells; and plant cells.

A great variety of expression systems can be used to produce a polypeptide
10 useful in the present invention. Such vectors include, among others,
chromosomal, episomal and virus-derived vectors, e.g., vectors derived from
bacterial plasmids, from bacteriophage, from transposons, from yeast episomes,
from insertion elements, from yeast chromosomal elements, from viruses such as
baculoviruses, papova viruses, such as SV40, vaccinia viruses, adenoviruses,
15 fowl pox viruses, pseudorabies viruses and retroviruses, and vectors derived
from combinations thereof, such as those derived from plasmid and
bacteriophage genetic elements, such as cosmids and phagemids. The
expression system constructs may contain control regions that regulate as well as
engender expression. Generally, any system or vector suitable to maintain.
20 propagate or express polynucleotides and/or to express a polypeptide in a host
may be used for expression in this regard. The appropriate DNA sequence may
be inserted into the expression system by any of a variety of well-known and
routine techniques, such as, for example, those set forth in Sambrook *et al.*

For secretion of the translated protein into the lumen of the endoplasmic
25 reticulum, into the periplasmic space or into the extracellular environment,
appropriate secretion signals may be incorporated into the expressed
polypeptide. These signals may be endogenous to the polypeptide or they may
be heterologous signals.

Polypeptides of the invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography is employed for purification. Well known techniques for refolding protein may be employed to regenerate active conformation when the polypeptide is denatured during isolation and/or purification.

10

Methods of Delivery

In the present invention the polynucleotide may be delivered to a target cell population, either *ex vivo* or *in vivo*, by any suitable Gene Delivery Vehicle.

15

This includes but is not restricted to, DNA, formulated in lipid or protein complexes or administered as naked DNA via injection or biolistic delivery, viruses such as retroviruses, adenoviruses, herpes viruses, vaccinia viruses, adeno associated viruses. The GDV can be designed by a person ordinarily skilled in the art of recombinant DNA technology and gene expression to express the fusion protein at appropriate levels and with the cellular specificity demanded by a particular application.

20

As it is well known in the art, a vector is a tool that allows or facilitates the transfer of an entity from one environment to another. In accordance with the present invention, and by way of example, some vectors used in recombinant DNA techniques allow entities, such as a segment of DNA (such as a heterologous DNA segment, such as a heterologous cDNA segment), to be transferred into a target cell. Optionally, once within the target cell, the vector may then serve to maintain the heterologous DNA within the cell or may act as a

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unit of DNA replication. Examples of vectors used in recombinant DNA techniques include plasmids, chromosomes, artificial chromosomes or viruses.

The vector can be delivered by viral or non-viral techniques.

5

Non-viral delivery systems include but are not limited to DNA transfection methods. Here, transfection includes a process using a non-viral vector to deliver a gene to a target mammalian cell.

- 10 Typical transfection methods include electroporation, DNA biolistics, lipid-mediated transfection, compacted DNA-mediated transfection, liposomes, immunoliposomes, lipofectin, cationic agent-mediated, cationic facial amphiphiles (CFAs) (Nature Biotechnology 1996 14: 556), multivalent cations such as spermine, cationic lipids or polylysine, 1, 2,-bis (oleoyloxy)-3-
15 (trimethylammonio) propane (DOTAP)-cholesterol complexes (Wolff and Trubetskoy 1998 Nature Biotechnology 16: 421) and combinations thereof.

- Viral delivery systems include but are not limited to adenovirus vector, an adeno-associated viral (AAV) vector, a herpes viral vector, a retroviral vector, a
20 lentiviral vector or a baculoviral vector.

- Examples of retroviruses include but are not limited to: murine leukemia virus (MLV), human immunodeficiency virus (HIV), equine infectious anaemia virus (EIAV), mouse mammary tumour virus (MMTV), Rous sarcoma virus (RSV),
25 Fujinami sarcoma virus (FuSV), Moloney murine leukemia virus (Mo-MLV), FBR murine osteosarcoma virus (FBR MSV), Moloney murine sarcoma virus (Mo-MSV), Abelson murine leukemia virus (A-MLV), Avian myelocytomatosis virus-29 (MC29), and Avian erythroblastosis virus (AEV).

A detailed list of retroviruses may be found in Coffin *et al* ("Retroviruses" 1997 Cold Spring Harbour Laboratory Press Eds: JM Coffin, SM Hughes, HE Varmus pp 758-763).

- 5 Adenoviruses and adeno-associated viruses which have good specificity for epithelial cells are particularly preferred.

Other examples of vectors include *ex vivo* delivery systems, which include but are not limited to DNA transfection methods such as electroporation, DNA
10 biolistics, lipid-mediated transfection, compacted DNA-mediated transfection.

Thus, nucleic acid vectors according to the invention may be capable of delivery preferentially to the target cell. For example in the case of a retroviral vector, the retroviral envelope protein may be capable of directing the vector to a
15 particular cell type or cell types. For that purpose, the envelope protein may be a modified envelope protein adapted to have a specific targeting ability, or it may be a selected envelope protein derived from a different viral or retroviral source and having the desired targeting ability.

20 Preferably, the nucleic acid in a vector according to the invention is operatively linked to an expression control sequence capable of causing preferential expression of the fusion protein in the target cell. The expression control sequence may be for example a promotor or enhancer which is preferentially active in certain cell types including the target cell, or a promotor or enhancer
25 which is preferentially active under certain conditions.

The term "promoter" is used in the normal sense of the art, e.g. an RNA polymerase binding site in the Jacob-Monod theory of gene expression.

The term "enhancer" includes a DNA sequence which binds to other protein components of the transcription initiation complex and thus facilitates the initiation of transcription directed by its associated promoter.

- 5 Preferably the promoters of the present invention are tissue specific. That is, they are capable of driving transcription of a nucleic acid in one tissue while remaining largely "silent" in other tissue types. A particularly preferred promoter is the epithelial cell promoter.
- 10 The term "tissue specific" means a promoter which is not restricted in activity to a single tissue type but which nevertheless shows selectivity in that they may be active in one group of tissues and less active or silent in another group.

Administration

- 15 Compounds capable of affecting a component of the Hedgehog family signalling pathway or a target pathway thereof for use in therapy are typically formulated for administration to patients with a pharmaceutically acceptable carrier or diluent to produce a pharmaceutical composition. The formulation will depend upon the nature of the compound identified and the route of administration but
- 20 typically they can be formulated for topical, parenteral, intramuscular, intravenous, intra-peritoneal, intranasal inhalation, lung inhalation, intradermal or intra-articular administration. The compound may be used in an injectable form. It may therefore be mixed with any vehicle which is pharmaceutically acceptable for an injectable formulation, preferably for a direct injection at the site to be
- 25 treated, although it may be administered systemically.

- The pharmaceutically acceptable carrier or diluent may be, for example, sterile isotonic saline solutions, or other isotonic solutions such as phosphate-buffered saline. The compounds of the present invention may be admixed with any
- 30 suitable binder(s), lubricant(s), suspending agent(s), coating agent(s),

solubilising agent(s). It is also preferred to formulate the compound in an orally active form.

5 In general, a therapeutically effective daily oral or intravenous dose of the compounds of the invention, including compounds of formula (1) and their salts, is likely to range from 0.01 to 50 mg/kg body weight of the subject to be treated, preferably 0.1 to 20 mg/kg. The compounds of the formula (I) and their salts may also be administered by intravenous infusion, at a dose which is likely to range from 0.001-10 mg/kg/hr.

10

Tablets or capsules of the compounds may be administered singly or two or more at a time, as appropriate. It is also possible to administer the compounds in sustained release formulations.

15 Typically, the physician will determine the actual dosage which will be most suitable for an individual patient and it will vary with the age, weight and response of the particular patient. The above dosages are exemplary of the average case. There can, of course, be individual instances where higher or lower dosage ranges are merited, and such are within the scope of this invention.

20

Alternatively, the compounds of the invention can be administered by inhalation or in the form of a suppository or pessary, or they may be applied topically in the form of a lotion, solution, cream, ointment or dusting powder. An alternative means of transdermal administration is by use of a skin patch. For example, they
25 can be incorporated into a cream consisting of an aqueous emulsion of polyethylene glycols or liquid paraffin. They can also be incorporated, at a concentration of between 1 and 10% by weight, into an ointment consisting of a white wax or white soft paraffin base together with such stabilisers and preservatives as may be required.

30

For some applications, preferably the compositions are administered orally in the form of tablets containing excipients such as starch or lactose, or in capsules or ovules either alone or in admixture with excipients, or in the form of elixirs, solutions or suspensions containing flavouring or colouring agents.

5

The compositions (as well as the compounds alone) can also be injected parenterally, for example intracavernosally, intravenously, intramuscularly or subcutaneously. In this case, the compositions will comprise a suitable carrier or diluent.

10

For parenteral administration, the compositions are best used in the form of a sterile aqueous solution which may contain other substances, for example enough salts or monosaccharides to make the solution isotonic with blood.

15 For buccal or sublingual administration the compositions may be administered in the form of tablets or lozenges which can be formulated in a conventional manner.

For oral, parenteral, buccal and sublingual administration to subjects (such as patients), the daily dosage level of the compounds of the present invention and their pharmaceutically acceptable salts and solvates may typically be from 10 to 500 mg (in single or divided doses). Thus, and by way of example, tablets or capsules may contain from 5 to 100 mg of active compound for administration singly, or two or more at a time, as appropriate. As indicated above, the physician will determine the actual dosage which will be most suitable for an individual patient and it will vary with the age, weight and response of the particular patient. It is to be noted that whilst the above-mentioned dosages are exemplary of the average case there can, of course, be individual instances where higher or lower dosage ranges are merited and such dose ranges are within
30 the scope of this invention.

The routes of administration and dosages described are intended only as a guide since a skilled practitioner will be able to determine readily the optimum route of administration and dosage for any particular patient depending on, for example,
5 the age, weight and condition of the patient.

The term treatment or therapy as used herein should be taken to encompass diagnostic and prophylactic applications.

10 The treatment of the present invention includes both human and veterinary applications.

As used herein the terms protein and polypeptide and peptide may be assumed to be synonymous, protein merely being used in a general sense to indicate a
15 relatively longer amino acid sequence than that present in a polypeptide, and polypeptide merely being used in a general sense to indicate a relatively longer amino acid sequence than that present in a peptide. Generally for ease of reference only we will simply refer to the term polypeptide.

20 The invention will now be described in further detail with reference to the following non-limiting examples:

Example 1 : Construction of SPC-Shh expression plasmid

25 The SPC-mouse Shh vector shown in Figure 5 was constructed from a parent expression vector that was made as follows. The parent vector shown in Figure 4 contains a pUC18 backbone with an ampicillin resistance gene, a 3.7 kb sequence containing the human SPC promoter region, a multiple cloning site (MCS), an SV40 small T intron and a 0.4 kb sequence containing a poly(A)
30 addition site and with stop codons in all three reading frame (Korfhagen *et al*;

Development 1997) .The cDNA sequence encoding the mouse Shh was cloned into the MCS.

Additional vectors including SPC-HIP, SPC-WIF-1, SPC-Dvl-1 were made by
5 cloning murine cDNAs for HIP, WIF-1, Dvl-1 into the MCS of the SPC expression vector.

**Example 2 : Increased expression of Ptc in the lung epithelial cells from human patients with idiopathic fibrosing alveolitis (IFA also known as
10 CFA) and in a murine model of interstitial pulmonary fibrosis (IPF).**

Fig. 6C

BALB/c mice were treated intratracheally with 50 µg of FITC dissolved in physiological buffered saline (PBS). Three months later mice were sacrificed
15 and lungs removed and fixed in 4% buffered formalin and embedded in paraffin. 5 µm sections of lung tissue were placed onto TESPA coated slides and the expression of Ptc gene expression was examined by anti-sense RNA in situ hybridization (ISH).

20 Sections were hybridized with digoxigenin antisense RNA probes specific for murine Ptc1 at 65°C . The bound probe was detected by alkaline phosphatase conjugated goat anti-digoxigenin Fab and sections were developed using NBT and BCIP as the substrate. We observed increased expression of Ptc in lung epithelial cells in the murine model of IPF. Expression of Ptc was restricted only
25 to those areas showing damage. Increased Ptc expression was noted within 24 hours of i.t. FITC and was maintained for at least 6 months.

Fig. 6 A & B

Paraffin embedded archive lung tissue from human patients diagnosed with IFA or control patients with healthy lung tissue were sectioned at 5 µm and placed onto TESPA coated slides. The slides were then analysed for Ptc gene expression by anti-sense RNA in situ hybridization (see above). We observed
5 increased expression of Ptc in lung epithelial cells.

Example 3 : Overexpression of Shh leads to epithelial cell hyperplasia and lung fibrosis. Figures 7-9

10 BALB/c mice were injected i.t. with either (i) saline alone, (ii) 20 µg of SPC plasmid dissolved in saline, or (iii) 20 µg of SPC-Shh plasmid DNA dissolved in saline on day 0 and day 5. The SPC plasmid provides tissue-specific expression of a desired gene as it contains the promoter sequence from the lung epithelial cell-specific surfactant protein C (i.e. SPC). Mice were sacrificed at day 12 and
15 day 35 where upon the lungs were removed and placed into 4% buffered formalin.

5 µm sections of lung tissue (Fig 7, 8) or trachea (Fig. 9) from each group at each of the two time points were placed onto poly-L-Lysine coated slides and
20 stained using the haematoxylin and eosin (H & E) histochemical stain.

The groups contained:

Day 12 PBS (2 mice), SPC (2 mice) and SPC-shh (2 mice)

Day 35 PBS (3 mice) SPC (3 mice) and SPC-shh (3 mice)

25 Slides from the day 35 treatment group were further analysed for collagen production using a Masons-trichrome histochemical stain. This type of stain is routinely used to stain for collagen fibres in tissue samples which turn green. Increased levels of collagen staining could be identified in lung tissue from SPC-Shh treated mice when compared to controls.

30

- Slides from the day 35 treatment group were also analysed for potential goblet cell hyperplasia using the periodic acid-schiff (PAS) histochemical stain. Using this stain, cells which generate mucins stain an intense pink colour while epithelium stains light blue. We observed an increase in the number of goblet cells in the lung tissue of SPC-Shh mice (i.e. pink cells) and increased mucous secretion into the airways. The increased goblet cell numbers cells were observed only in those airways showing signs of epithelia hyperplasia. Normal airways in the SPC-Shh mice were equivalent to those in the control mice.
- 10 Slides from Day 12 and Day 35 treatment groups were stained for a proliferation index marker, Ki-67, to provide evidence that the lung epithelia of SPC-shh mice were actively proliferating. Slides from the three treatment groups were stained with an antibody specific for the Ki-67 antigen which marks cells in the S-phase of the cell cycle. There was an increase in the number of Ki-67 +ve cells in the lung epithelium of SPC-Shh treated mice when compared to the epithelium of control mice
- 15

Example 4 : Epithelial cells express high levels of Shh following FITC damage

- 20 Mice were treated intratracheally with the hapten fluorescein isothiocyanate. Seven days later the lungs were removed and fixed in formalin. Sections were cut and stained for Shh by immunohistochemistry. Figs. 10A and 10B show expression of Shh in the lung of FITC treated mice, while Fig. 10C shows the staining for Shh observed in the control lung. Shh could be detected on epithelial cells, and a higher level of Shh was detected on a basal cell population in the lung interstitium consistent in morphology with fibroblasts.
- 25

Example 5 : Epithelial cells express high levels of Shh following FITC damage

5 Mice were treated intratracheally with the hapten fluorescein isothiocyanate. One month later the lungs were removed and fixed in formalin. Sections were cut and stained for Shh by immunohistochemistry. Figs. 11A and 11B show expression of Shh in two different sections of lung of FITC treated mice, while Figs. 11C and 11D show the staining for Shh observed in the control lung. Shh
10 could be detected at a higher level on epithelial cells than on a basal cell population in the lung interstitium.

Example 6 : Epithelial cells express high levels of Ptc following FITC damage

15 Mice were treated intratracheally with the hapten fluorescein isothiocyanate. Seven days later the lungs were removed and fixed in formalin. Sections were cut and stained for Shh by immunohistochemistry. Fig. 12A shows expression of Ptc in the lung of FITC treated mice, while Fig. 12B shows the staining for Ptc
20 observed in the control lung. Ptc could be detected on epithelial cells and a higher level of Ptc was detected on infiltrating leukocytes found in the lung interstitium.

Example 7 : Shh and Ptc staining on biopsy material from human CFA lung

25 Archive material from a CFA patient was sectioned and stained by immunohistochemistry for the presence of Shh-N (the bioactive protein) and Ptc. Figs. 13A-C show staining for Shh and Figs. 13D-F show Ptc expression. Each Fig. represents a serial section taken from the same piece of lung at 10x. The

cells within the airway interstitium and aveolar space contain leukocytes and these stain strongly for Ptc.

Example 8 : Shh and Ptc staining on biopsy material from human CFA lung

5

Archive material from a CFA patient was sectioned and stained by immunohistochemistry for the presence of Shh-N (the bioactive protein) and Ptc. Figs. 14A and B show staining for Shh and Figs. 14C and D show Ptc expression. Figs. 14A and C represents a serial section taken from the same
10 piece of lung at 10x, while Figs. 14B and D are 40x views taken from the lower portion of the section. The cells within the airway interstitium and aveolar space contain leukocytes and these stain strongly for Ptc.

Example 9 : Shh and Ptc staining on biopsy material from human CFA lung

15

Archive material from a CFA patient was sectioned and stained by immunohistochemistry for the presence of Shh-N (the bioactive protein) and Ptc. Figs. 15A and B show staining for Shh and Figs. 15C and D show Ptc expression. Figs. 15A and C represent serial sections taken from the same
20 of lung, while Figs. 15B and D are serial sections taken from another section. The cells within the airway lumen contain aveolar macrophages and these stain strongly for Ptc.

Example 10 : Effect of introduction of SPC-HIP

25

Our previous studies have revealed that dysregulation of the Shh signalling pathway during epithelial cell repair in the lung, can lead to lymphocyte infiltration with concomitant induction of interstitial fibrosis and scarring. We have previously used a novel model of pulmonary fibrosis where naive BALB/c
30 mice were treated i.t. with 50 µg of FITC dissolved in saline (PBS) leads to an

initial strong inflammatory response which resolves by day 7, but EC hyperplasia is evident at this time and lymphocytes begin to infiltrate the lung at the sites of EC hyperplasia by day 21. By day 28 there is evidence of interstitial fibrosis which seems to be aggravated by the presence of lymphocytes in the lung. In this model we have observed increased expression of Ptc-1 in sites of EC hyperplasia but not in normal areas of lung tissue. This indicates that there is dysregulation of the Shh pathway in the disease process in the FITC treated mice.

10 Since HIP is a natural antagonist of the Shh protein, we overexpress HIP in the lung using the SPC expression vector as follows:

BALB/c mice are treated i.t. with 50 µg of FITC dissolved in saline (PBS) and 1-3 months later mice, which are time points where it is known that mice would normally have mild to severe fibrosis respectively, they are given two injections of either SPC plasmid alone in saline or SPC-HIP in saline i.t. 7 days apart. Mice are sacrificed at day 7, day 30 and day 60 post SPC-HIP administration. The lung tissue is removed and fixed in 4% buffered formalin. Sections are examined by H & E, PAS, Maser's-trichrome and Ki-67 at the various time points. In addition, we examine the expression of Ptc and Shh by ISH and immunohistochemistry. A reduction of Ptc expression and EC hyperplasia and lung fibrosis is seen when compared to the epithelium of control mice.

Example 11 : Effect of introduction of PKA

25 BALB/c mice are treated i.t. with 50 µg of FITC dissolved in saline (PBS) and 1-3 months later mice, which are time points where it is known that mice would normally have mild to severe fibrosis respectively, they are given two injections of PKA in saline i.t. 7 days apart. Mice are sacrificed at day 7, day 30 and day 30 60 post SPC-HIP administration. The lung tissue is removed and fixed in 4%

buffered formalin. Sections are examined by H & E, PAS, Masons-trichrome and Ki-67 at the various time points. In addition, we examine the expression of Ptc and Shh by ISH and immunohistochemistry. A reduction of Ptc expression and EC hyperplasia and lung fibrosis is seen when compared to the epithelium of control mice.

Example 12 : Epithelial cells express high level of Dvl-1 following FITC damage

Mice were treated intratracheally with the hapten fluorescein isothiocyanate. Seven days later the lungs were removed and fixed in formalin. Sections were cut and stained for Dvl-1 by immunohistochemistry. Fig. 16A shows expression of Dvl-1 in the lung of FITC treated mice, while Fig. 16B shows the staining for Dvl-1 observed in the control lung. Strong expression of Dvl-1 on epithelial cells and infiltrating leukocytes was observed.

Example 13 : Dvl-1 adenovirus induces epithelial cell proliferation

Mice were treated intratracheally with a control adenovirus (Figs. 17A and C) or an adenovirus containing the murine Dvl-1 cDNA (Figs. 17B and D). Four days later the lungs were removed and fixed in formalin. Sections were cut and stained for the proliferation marker Ki67 by immunohistochemistry. Figs. 17A and B show a view of the lung at 10x and Figs. 17C and D are 40x views of the sections shown in the dotted box. Proliferating cells express the Ki67 antigen at high levels.

Example 14 : A role for dysregulation of Wnt signalling in lung fibrosis

The Dvl-1 protein is overexpressed in the lung epithelia of mice to examine what effect it may have on the development of lung fibrosis.

BALB/c mice were injected i.t. with either (i) 20 µg of SPC plasmid dissolved in saline, or (ii) 20 µg of SPC-Dvl-1 plasmid DNA dissolved in saline on day 0 and day 7. Mice were sacrificed at day 14 and day 35 where upon the lungs were removed and placed into 4% buffered formalin.

5

The lung tissue is removed and fixed in 4% buffered formalin. Sections are examined by H & E, PAS, Masons-trichrome and Ki-67 at the various time points. In addition, the expression of Dvl-1, Ptc and Shh is examined by ISH and immunohistochemistry. It was found that overexpression leads to EC

10 hyperplasia and lung fibrosis.

Example 15 : To examine the effect of introduction of the Wnt antagonist WIF-1

- 5 WIF-1 is a novel protein that was recently identified to be expressed as a transmembrane protein which binds to Wnt proteins to neutralize them. WIF-1 is normally expressed in the lung tissue, as well as the brain and so we performed a similar series of experiments as for the SPC-HIP protocols using SPC-WIF-1 in its place. Again a reduction in EC hyperplasia and lung fibrosis is seen when compared to the epithelium of control mice.

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Claims

1. Use of an inhibitor of a Hedgehog signalling pathway, or an inhibitor of a
5 pathway which is a target of the Hedgehog signalling pathway in the preparation
of a medicament for treatment of epithelial cell hyperplasia, fibrosis of tissue,
inflammation, cancer or an immune disorder.
2. Use of claim 1 wherein the Hedgehog signalling pathway is the Sonic
10 hedgehog, Indian hedgehog or Desert hedgehog signalling pathway.
3. Use of claim 1 wherein the pathway which is a target of the Hedgehog
signalling pathway is the Wnt or BMP signalling pathway.
- 15 4. Use of any preceding claim in which the inhibitor is HIP, cyclopamine,
Fzb, Cerberus, WIF-1, Xnr-3, Noggin, Chordin, Gremlin, or Follistatin or a
derivative, fragment, variant, mimetic, homologue or analogue thereof.
5. Use of any one of claims 1 to 4 in which the inhibitor is Ptc, Cos2 or
20 PKA or an agent of the cAMP signal transduction pathway.
6. Use of any one of claims 1 to 4 wherein the inhibitor is an antibody.
7. Use of any preceding claim for the preparation of a medicament for the
25 treatment of the lung or kidney.
8. Use of any preceding claim for the preparation of a medicament for the
treatment of adult respiratory distress syndrome; chronic obstructive airway
disorders including asthma, emphysema and chronic bronchitis; atelectasis;

occupational lung disease including silicosis; hypersensitivity diseases of the lung including

hypersensitivity pneumonitis; idiopathic interstitial lung diseases including
5 idiopathic pulmonary fibrosis, pneumonia including usual interstitial pneumonia, desquamative interstitial pneumonia and acute interstitial pneumonia; and pleural fibrosis.

9. Use of any preceding claim in which the immune disorder is an
10 autoimmune disease or graft rejection.

10. Use according to claim 9 in which the autoimmune disease is thyroiditis, insultitis, multiple sclerosis, iridocyclitis, uveitis, orchitis, hepatitis, Addison's disease, myasthenia gravis, rheumatoid arthritis and lupus erythematosus.

15

11. Use according to any preceding claim in which the cancer is an adenocarcinoma.

12. A composition for use in treatment of epithelial cell hyperplasia, fibrosis
20 of tissue, inflammation, cancer or an immune disorder comprising a therapeutically effective amount of an inhibitor of a Hedgehog signalling pathway or an inhibitor of a target pathway of the Hedgehog signalling pathway and a pharmaceutically acceptable carrier, diluent or excipient.

25 13. A method for identifying a compound that is an inhibitor of a Hedgehog signalling pathway or a target pathway of the Hedgehog signalling pathway comprising the steps of: (a) determining the activity of the signalling pathway in the presence and absence of said compound; (b) comparing the activities observed in step (a); and (c) identifying said compound as inhibitor by the

observed difference in the activity of the pathway in the presence and absence of said compound.

14. Use of an inhibitor identifiable using the method of claim 13 in a use of
5 any one of claims 1 to 11 or a composition of claim 12.

15. A vector capable of expressing an inhibitor of a Hedgehog signalling pathway or a target pathway of the Hedgehog signalling pathway.

10 16. A transgenic animal or cell line capable of expressing an inhibitor of a Hedgehog signalling pathway or a target pathway of the Hedgehog signalling pathway.

17. A transgenic animal or cell line according to claim 16 wherein the
15 inhibitor is WIF-1, Fzb-1, Noggin or HIP.

18. A transgenic animal or cell line capable of expressing a component of the Hedgehog signalling pathway or a component of pathway which is a target of the Hedgehog signalling pathway.

20 19. A transgenic animal or cell line according to claim 18 wherein the component is Sonic Hedgehog.

20. Use of a transgenic animal or cell line according to any of claims 16 to
25 19 as a disease model or in a method according to claim 13.

1 / 15

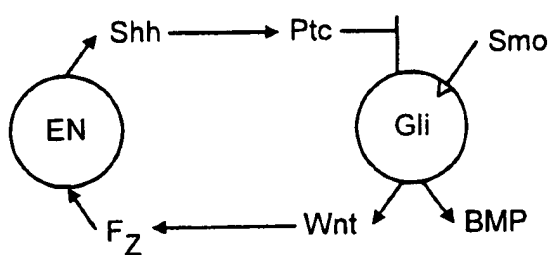


FIG. 1

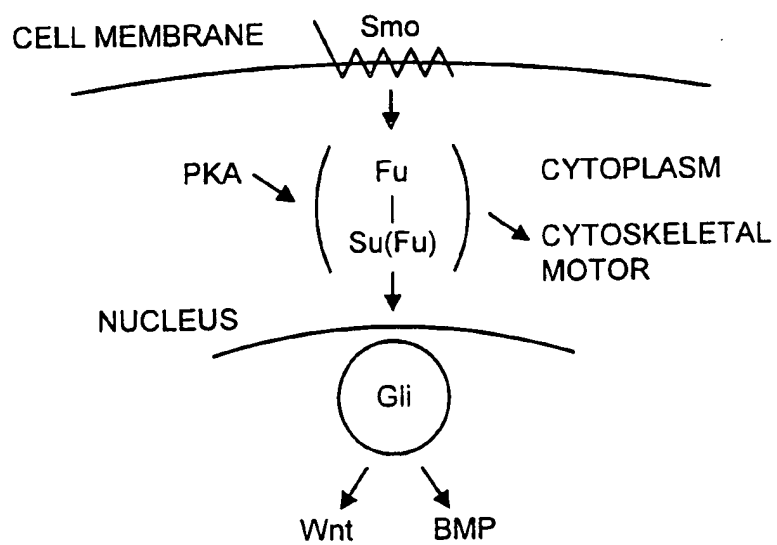


FIG. 2

2 / 15

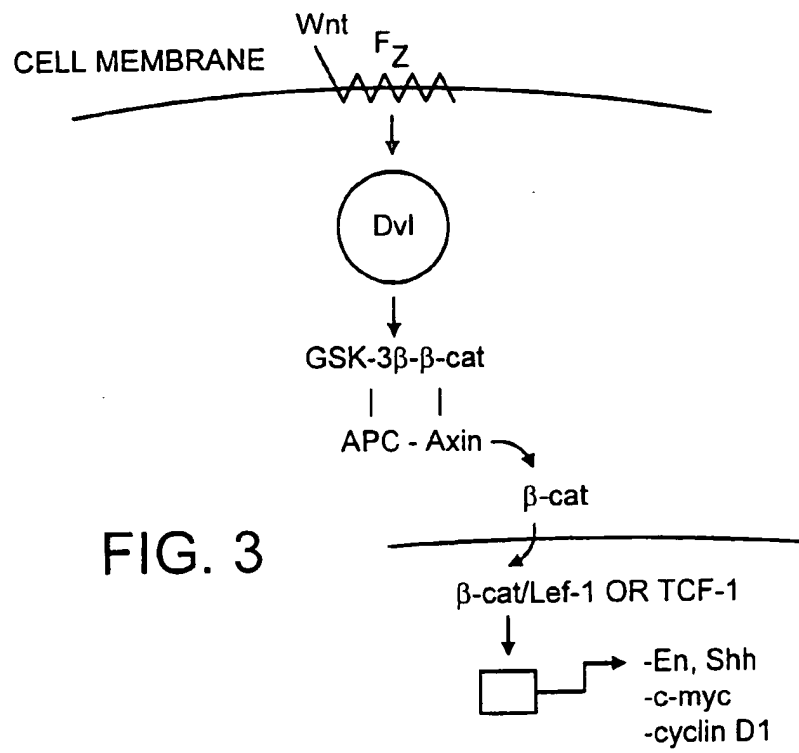


FIG. 3

3 / 15

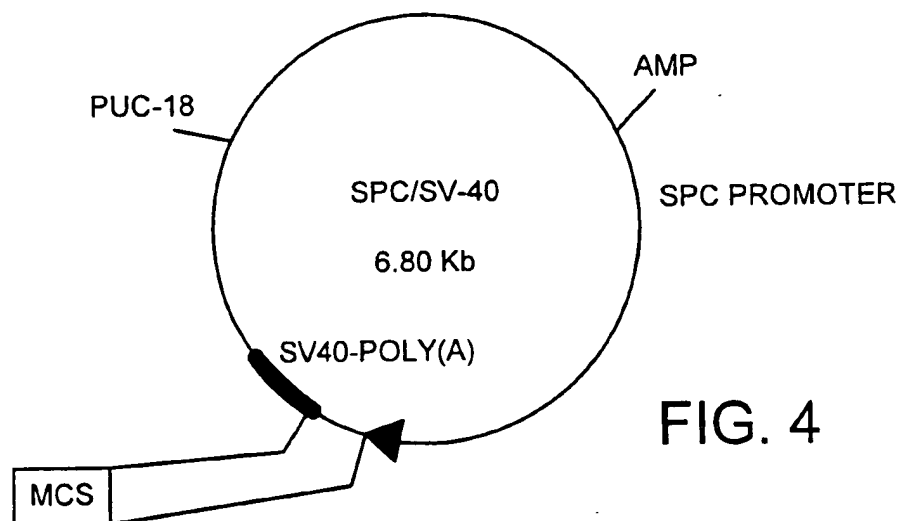


FIG. 4

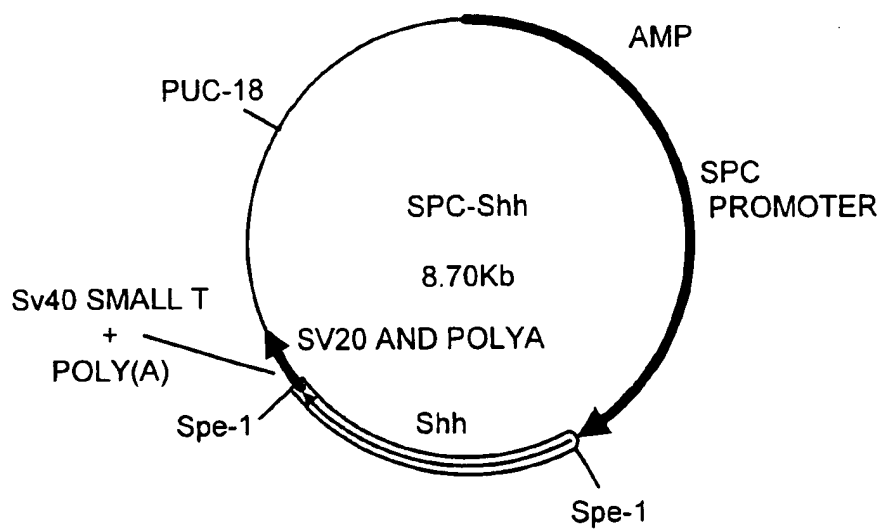
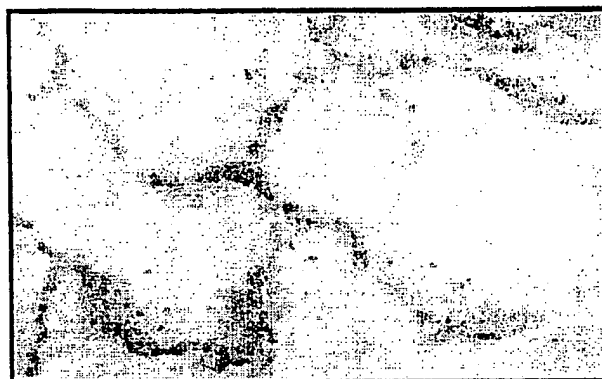


FIG. 5

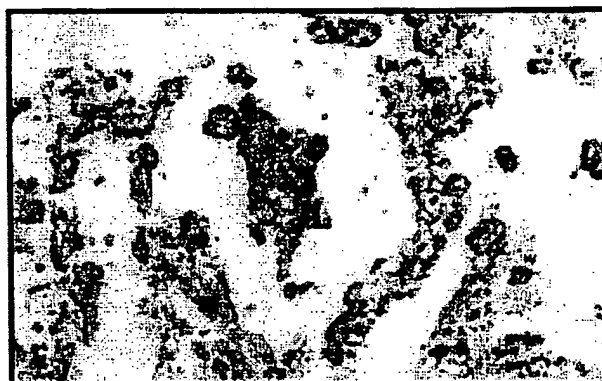
4 / 15

FIG. 6

PTC *in situ* HYBRIDISATION



a) NORMAL HUMAN LUNG



b) CFA LUNG



c) MOUSE LUNG 3 WEEKS POST FITC

SUBSTITUTE SHEET (RULE 26)

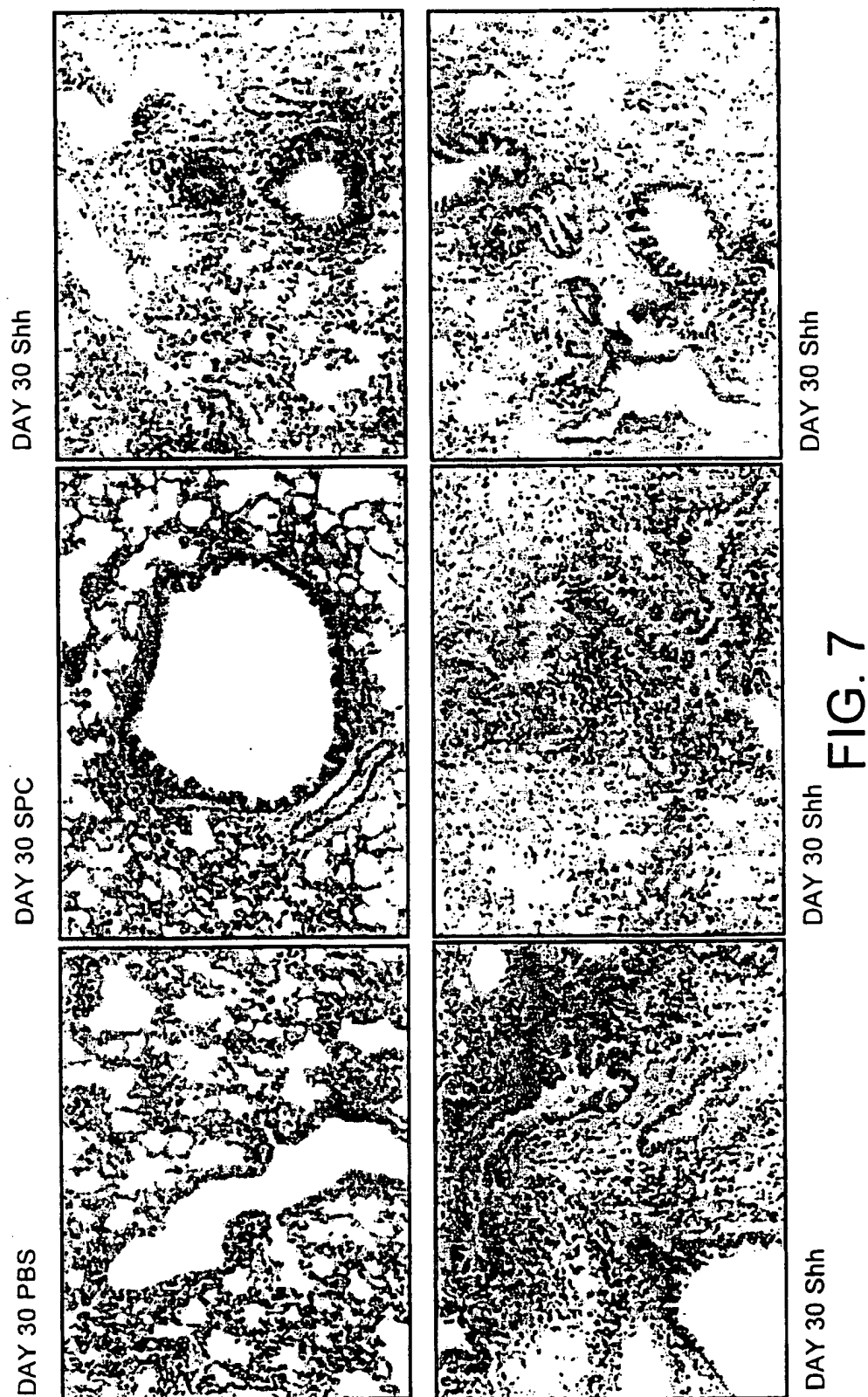
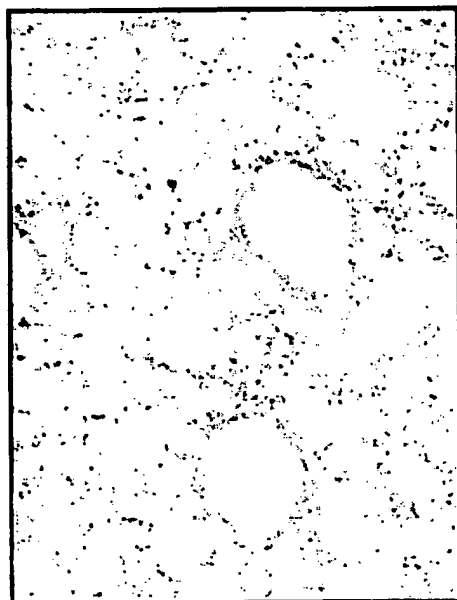


FIG. 8

Shh INDUCED EC HYPERPLASIA

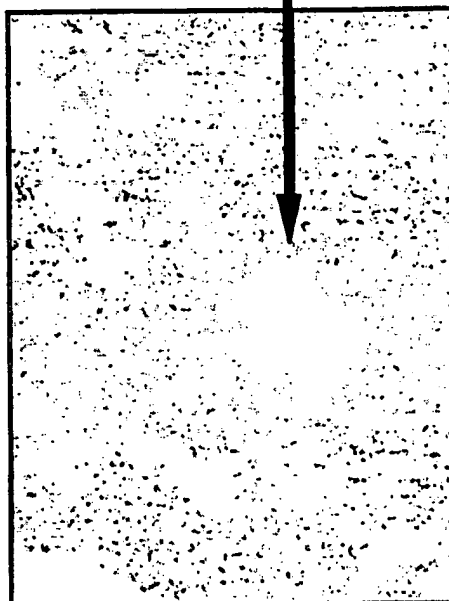
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DAY 7 SPC



DAY 7 Shh



EC HYPERPLASIA



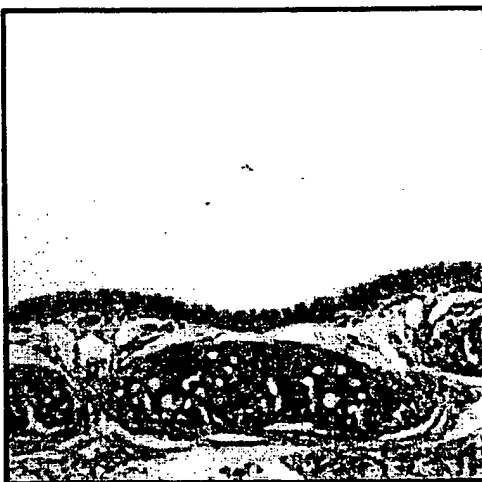
7 / 15

FIG. 9
TRACHEA

DAY 30 Shh



DAY 30 SPC



DAY 30 PBS

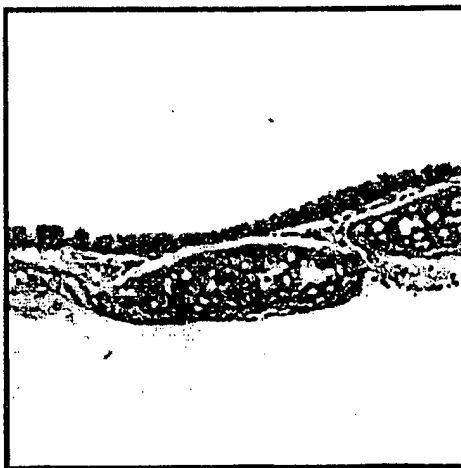
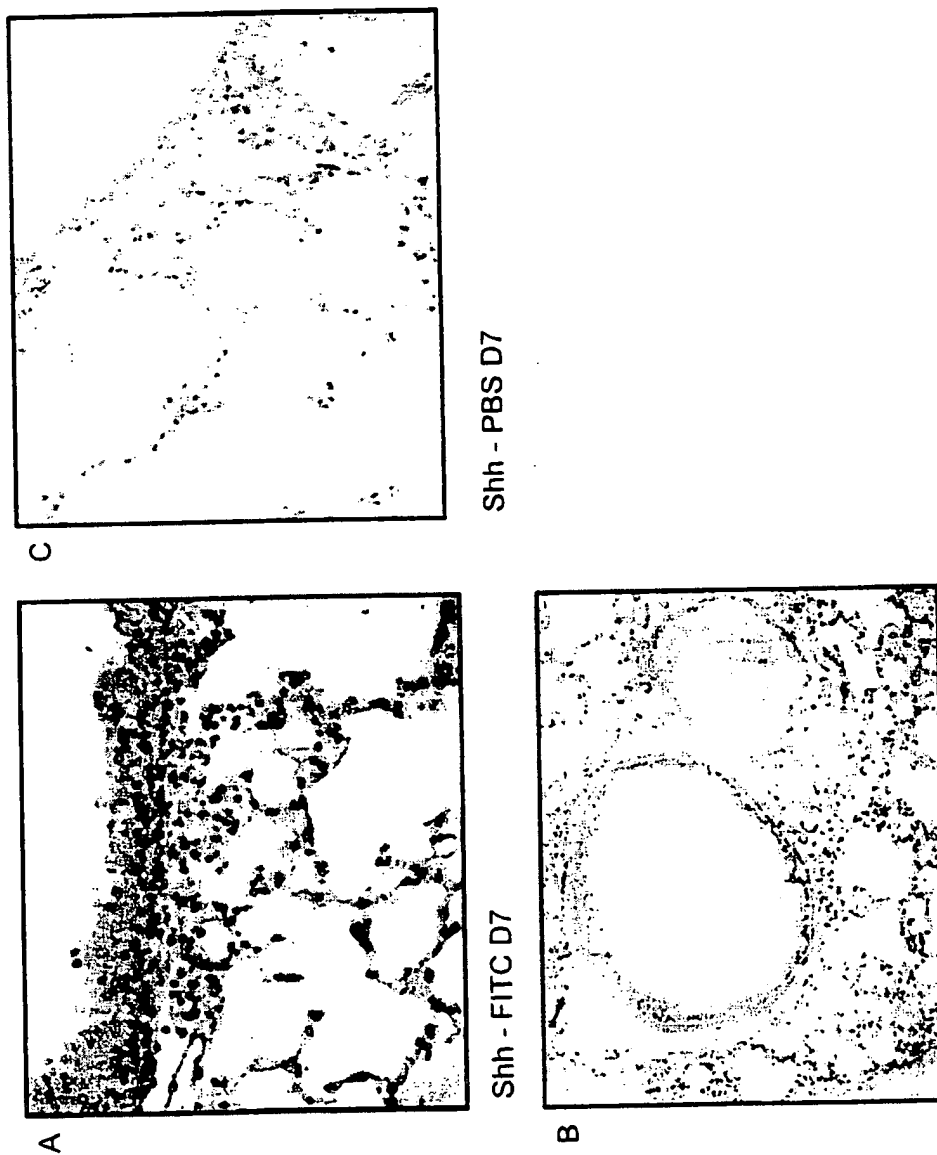


FIG. 10

Shh STAINING OF FITC LUNGS



9 / 15

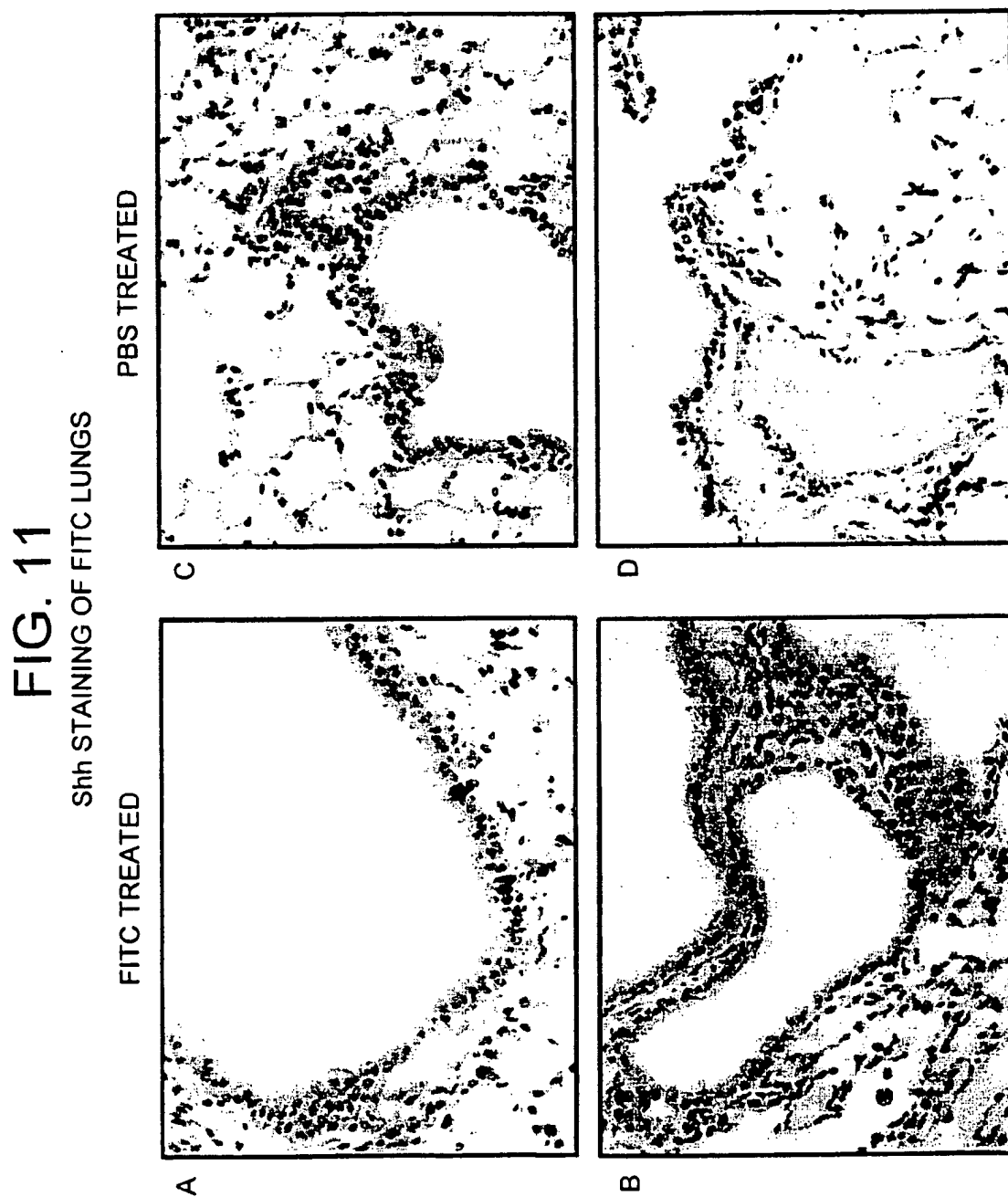


FIG. 12
Ptc STAINING OF FITC LUNGS

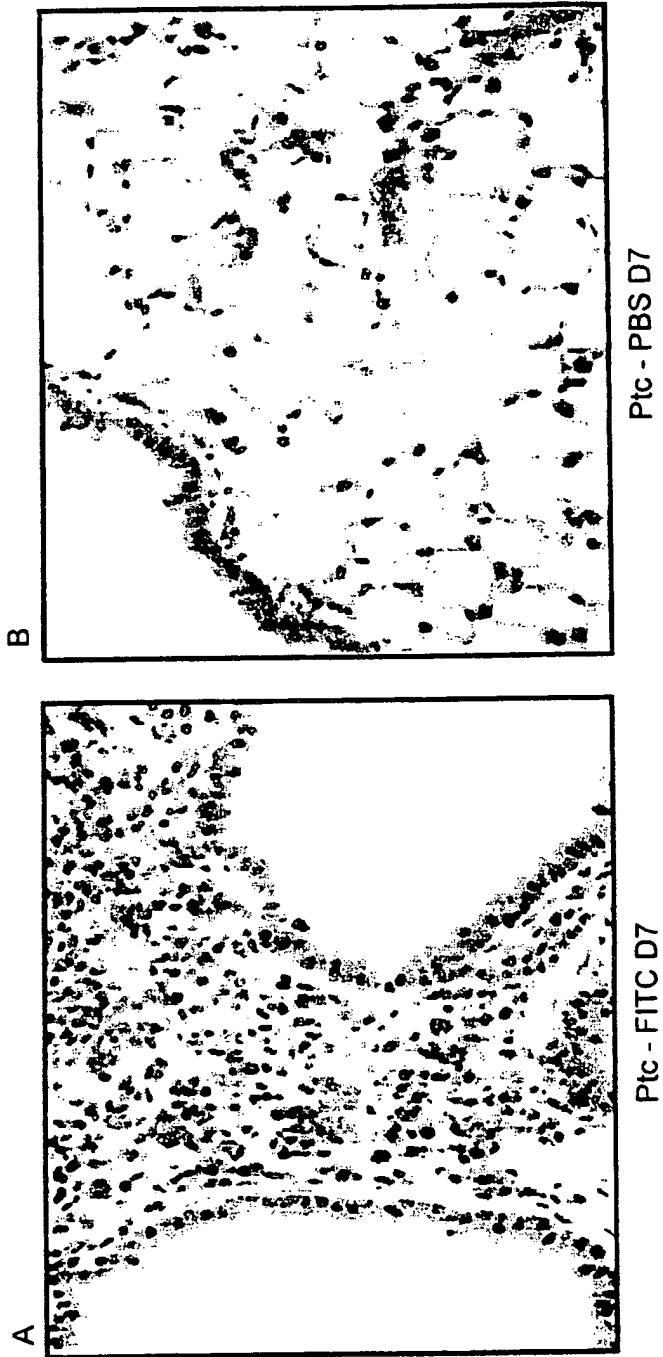


FIG. 13

CFA PATIENT

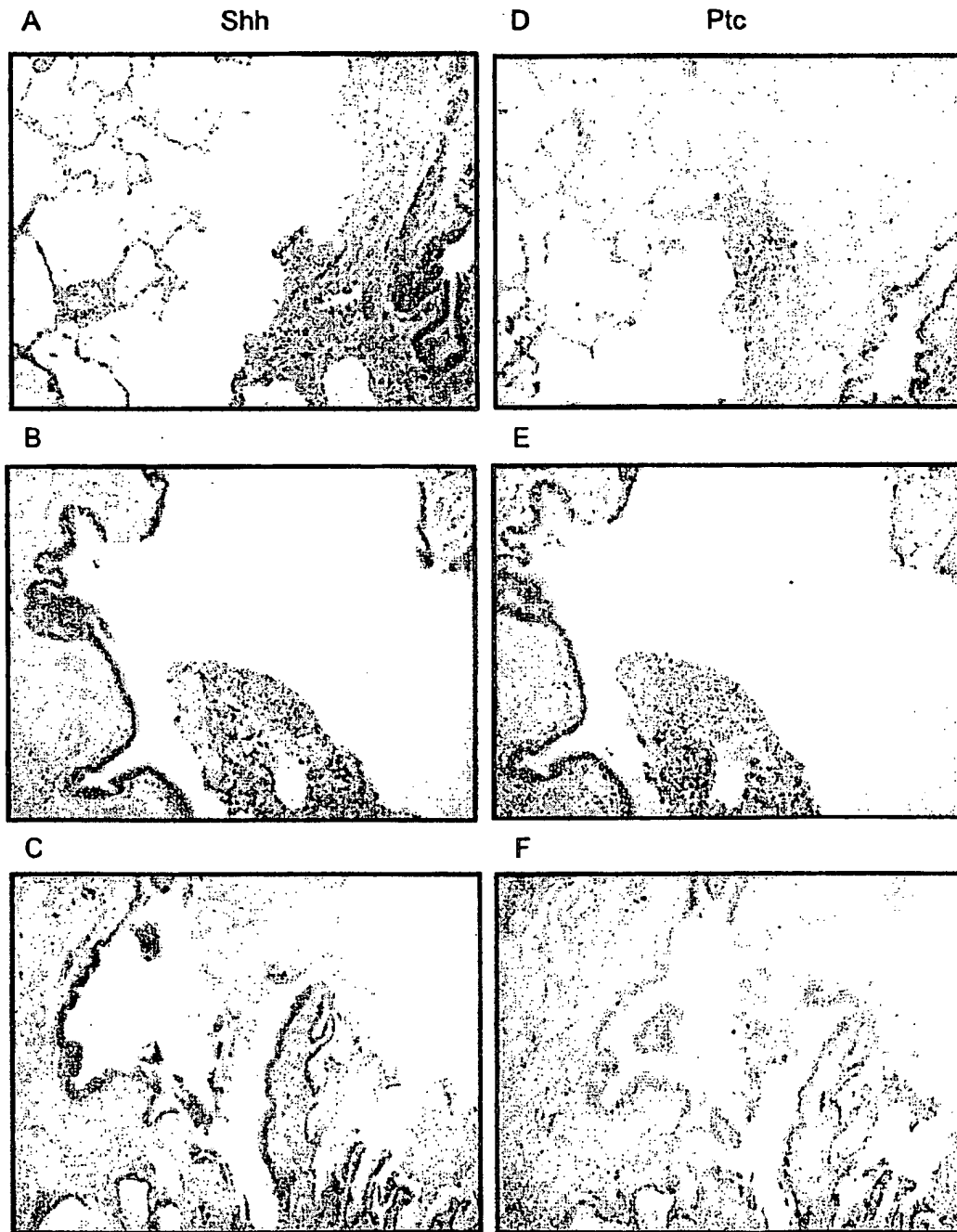


FIG. 14
CFA PATIENT

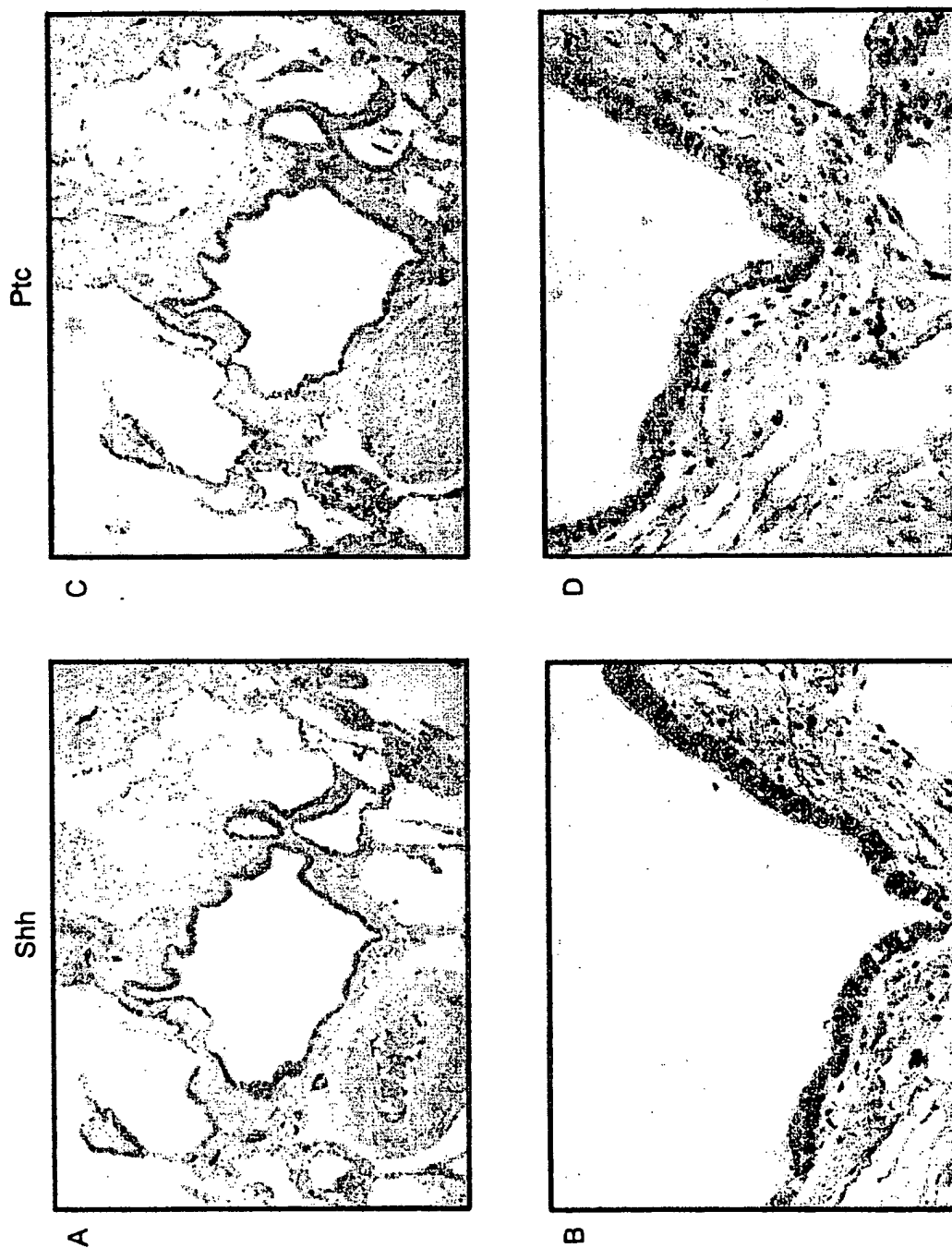


FIG. 15
CFA PATIENT

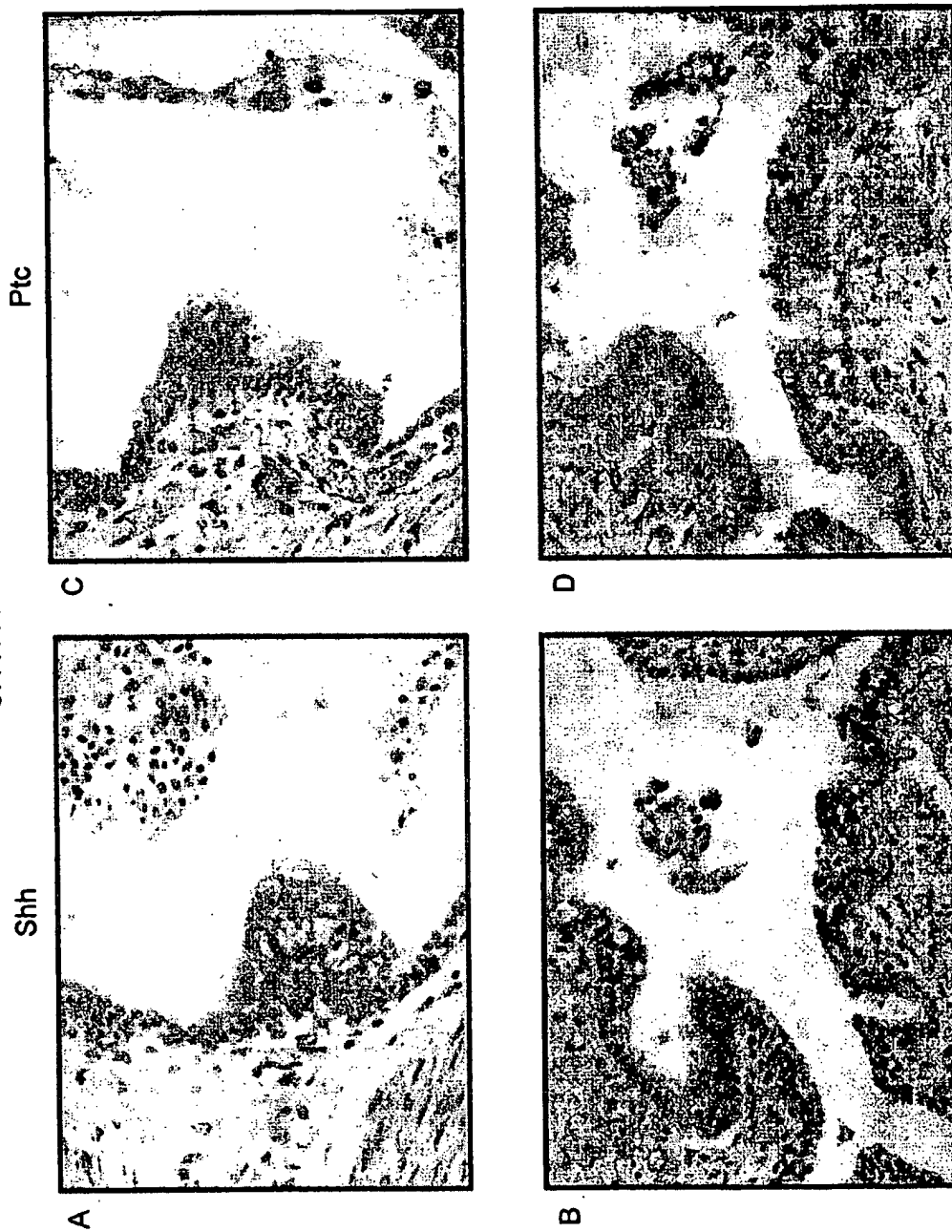


FIG. 16
DVL-1 STAINING OF FITC LUNGS

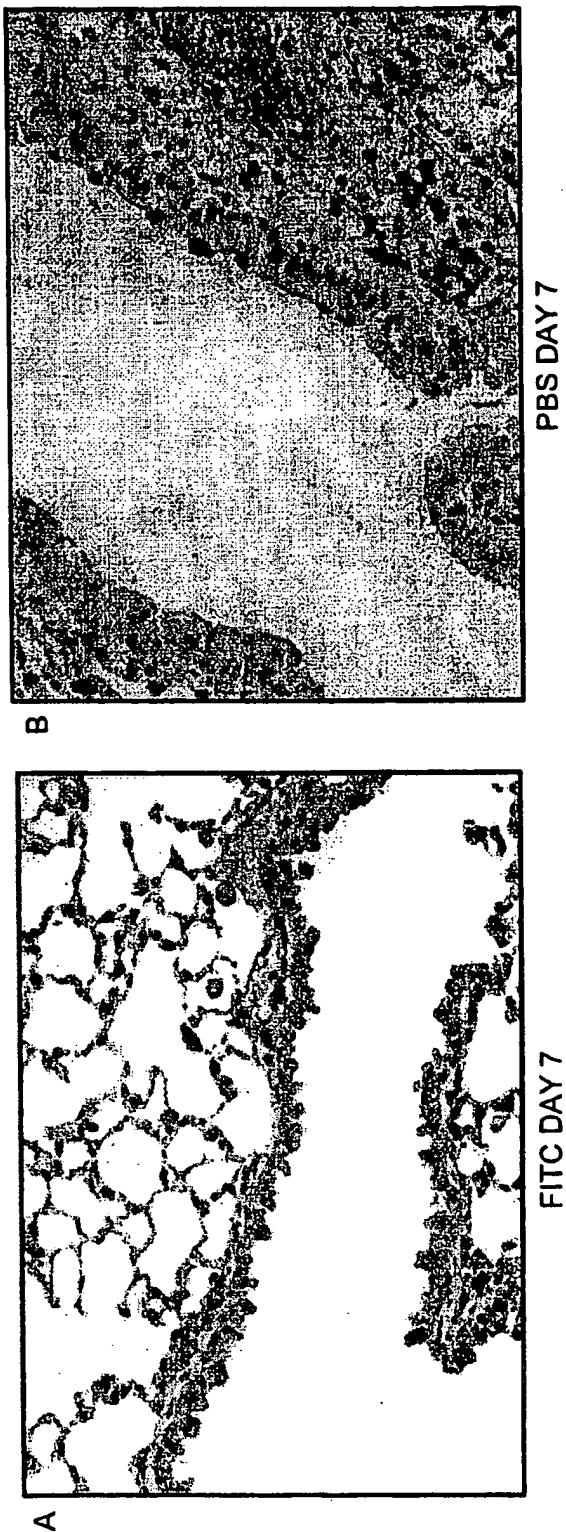
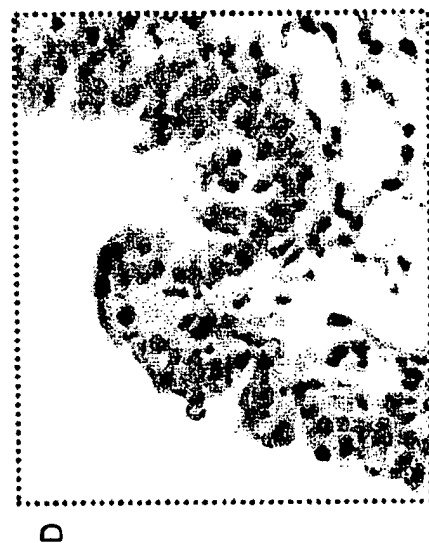
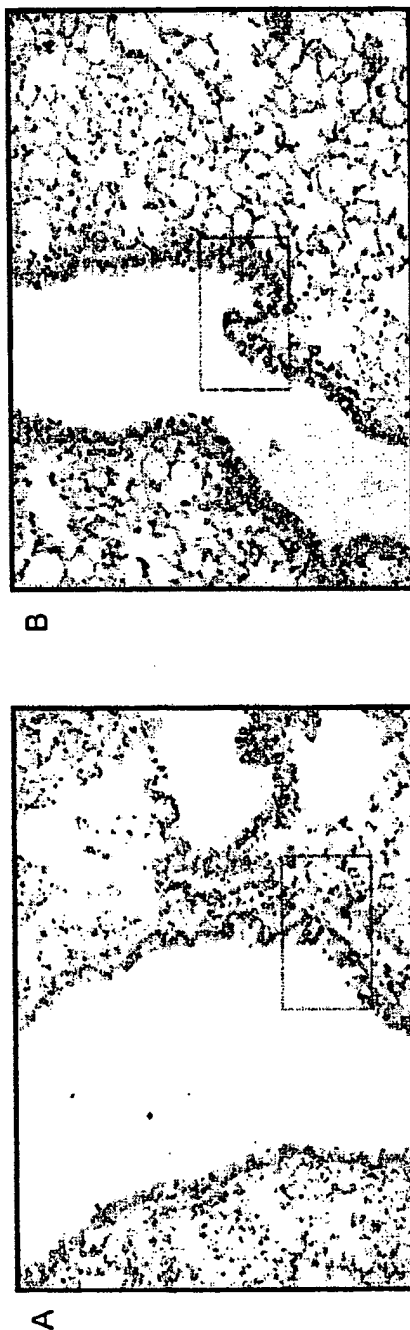
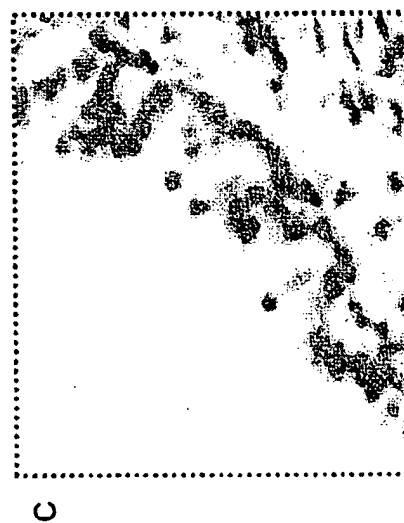


FIG. 17

DVL 1 ADVENOVIRUS *in vivo* (Ki67)



DAY 4 DVL-1



DAY 4 CONTROL VIRUS

SEQ ID NOS: 1 & 2

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 SOURCE house mouse.
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SEQ ID NOS: 3 & 4

DEFINITION Mus musculus C57 Black/6 dishevelled segment polarity protein homolog (Dvl-1) mRNA, complete cds.

ACCESSION U10115

NID g497689

KEYWORDS .

SOURCE mouse.

ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
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FEATURES Location/Qualifiers

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SEQ ID NOS: 5 & 6

DEFINITION Mus musculus hedgehog-interacting protein (Hip) mRNA, complete cds.

ACCESSION AF116865

NID g4868121

KEYWORDS .

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

FEATURES Location/Qualifiers

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BASE COUNT 660 a 722 c 672 g 615 t
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SEQ ID NOS: 7 & 8

DEFINITION Mus musculus Wnt inhibitory factor-1 mRNA, complete cds.

ACCESSION AF122923

NID g4585371

KEYWORDS .

SOURCE house mouse.

ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

FEATURES Location/Qualifiers

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INTERNATIONAL SEARCH REPORT

national Application No PCT/GB 00/02191					
A. CLASSIFICATION OF SUBJECT MATTER IPC 7 A61K38/17 A61K39/395 A61K48/00 G01N33/53 C12N15/63 A01K67/027 A61P11/00 A61P13/12 A61P35/00 A61P37/00					
According to International Patent Classification (IPC) or to both national classification and IPC					
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 C07K					
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched					
Electronic data base consulted during the International search (name of data base and, where practical, search terms used) BIOSIS, EMBASE, WPI Data, PAJ, EPO-Internal					
C. DOCUMENTS CONSIDERED TO BE RELEVANT					
Category *	Citation of document, with indication, where appropriate, of the relevant passages				Relevant to claim No.
X	J. DUMONTELLE ET AL.: "Inhibition of expression of the endogenous mammary oncogene Wnt-1 by antisense oligodeoxynucleotides." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH, vol. 36, March 1995 (1995-03), page 515 XP002147584 USA abstract 3065 <div style="text-align: center; margin-top: 20px;">--- -/--</div>				1,3, 12-14
<div style="display: flex; justify-content: space-between;"> <div> <input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. </div> <div> <input checked="" type="checkbox"/> Patent family members are listed in annex. </div> </div>					
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 50%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"G" document member of the same patent family</p> </div> </div>					
Date of the actual completion of the international search <div style="text-align: center; font-weight: bold;">15 September 2000</div>				Date of mailing of the international search report <div style="text-align: center; font-weight: bold;">04/10/2000</div>	
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3018				Authorized officer <div style="text-align: center; font-weight: bold;">Nooij, F</div>	

INTERNATIONAL SEARCH REPORT

national Application No
PCT/GB 00/02191

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	E. FUJITA ET AL.: "Involvement of sonic hedgehog in the cell growth of LK-2 cells, human lung squamous carcinoma cells." BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, vol. 238, no. 2, 18 September 1997 (1997-09-18), pages 658-664, XP002147585 Duluth, MN, USA abstract figure 5	1,2,6,7, 12-14
X	WO 99 22000 A (DEUTSCHES KREBSFORSCHUNGSZENTRUM STIFTUNG DES ÖFFENTLICHEN RECHTS) 6 May 1999 (1999-05-06) the whole document	1,3,6,9, 12-15
X	M. MURONE ET AL.: "Sonic hedgehog signaling by the patched-smoothened receptor complex." CURRENT BIOLOGY, vol. 9, no. 2, 28 January 1999 (1999-01-28), pages 76-84, XP000929784 the whole document	1-3,5, 13-15
X	J. POLANEC ET AL.: "Effect of Wnt-1 antisense RNA on the outgrowth of a mammary adenocarcinoma cell line expressing that oncogene." CLINICAL MOLECULAR PATHOLOGY, vol. 49, no. 3, 1996, pages M166-M169, XP000929749 abstract	1,3, 12-16
A	EP 0 874 048 A (KABUSHIKI KAISHA HAYASHIBARA SEIBUTSU KAGAKU KENKYUJO) 28 October 1998 (1998-10-28) examples 4,5 claims	1-20
P,X	WO 99 52534 A (JOHNS HOPKINS UNIVERSITY SCHOOL OF MEDICINE) 21 October 1999 (1999-10-21) examples claims	1,2,7,8, 12-14
P,X	WO 99 63052 A (THE PRESIDENT AND FELLOWS OF HARVARD COLLEGE) 9 December 1999 (1999-12-09) examples claims	18-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

national Application No

PCT/GB 00/02191

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
W0 9922000	A	06-05-1999	DE 19747418 C EP 1027440 A	15-07-1999 16-08-2000
EP 874048	A	28-10-1998	JP 11004695 A	12-01-1999
W0 9952534	A	21-10-1999	AU 3486099 A AU 3479099 A	01-11-1999 01-11-1999
W0 9963052	A	09-12-1999	AU 4416499 A	20-12-1999